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(21) International Application Number: PCT/US97/02930 (22) International Filing Date: 20 February 1997 (20.02.97) (30) Priority Data: 08/603,737 20 February 1996 (20.02.96) US (60) Parent Application or Grant (63) Related by Continuation US 08/603,737 (CIP) Filed on 20 February 1996 (20.02.96) (71) Applicant (for all designated States except US): THE GOVERNMENT OF THE UNITED STATES OF AMERICA, represented by THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES [US/US]; Office of Technology Transfer, National Institutes of Health, Suite 325, 6011 Executive Boulevard, Rockville, MA 20852 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): MAJER, Pavel [CZ/US]; 785H Wembley Drive, Frederick, MA 21701 (US). COLLINS, Jack [US/US]; 157 Stonegate Drive, Frederick, MA 21702 (US). GULNIK, Sergei [RU/US]; 8004 Meadowview Drive, Frederick, MA 21702 (US). ERICKSON,	John [US/US]; 5406 Jefferson Boulevard, Frederick, MA 21701 (US). PAN, Wenxi [CN/US]; 790B Wembley Drive, Frederick, MA 21701 (US). EISSENSTAT, Michael, Allen [US/US]; 8110 Claiborne Court, Frederick, MA 21702 (US). (74) Agents: FEILER, William, S. et al.; Morgan & Finnegan, L.L.P., 345 Park Avenue, New York, NY 10154 (US). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU. ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(54) Title: PEPTIDOMIMETIC INHIBITORS OF CATHEPSIN D AND PLASMEPSINS I AND II (57) Abstract <p>The present invention relates to the design and synthesis of linear and cyclic inhibitors of cathepsin D and plasmepsins I and II. The present invention also relates to the uses of these inhibitors for inhibiting invasion and metastasis of cancerous cells. The present invention further relates to the use of cathepsin D and plasmepsin I and II inhibitors for the prevention and treatment of Alzheimer's disease and malaria.</p>		

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PEPTIDOMIMETIC INHIBITORS OF CATHEPSIN D AND PLASMEPSINS I AND II

FIELD OF THE INVENTION

The invention relates to linear and cyclic inhibitor compounds of cathepsin D and plasmepsins I and II, and the use of these compounds for the prevention and treatment of diseases, and the like.

BACKGROUND OF THE INVENTION

Human cathepsin D is an intracellular aspartic protease normally found in lysosomes of all cells. The main physiological action of cathepsin D is degradation of cellular and phagocytosed proteins. Cathepsin D has also been implicated in a number of diseases. Elevated levels of cathepsin D have been correlated with poor prognosis in breast cancer. It has also been correlated with increase cell invasion and increased risk of metastasis, as well as shorter relapse-free survival. (See Rochefort, H., Semin. Cancer Biol. 1: 153 (1990) and Tandon, A. K. et al. N. Eng. J. Med. 322, 297 (1990)). The increased level of secretion of cathepsin D in breast cancer cells is due to both overexpression of the gene and altered processing of the protein. High levels of cathepsin D and other proteases, such as collagenase, produced in the vicinity of the growing tumor may degrade the extracellular matrix and thereby promote the escape of cancer cells to the lymphatic and circulatory systems and enhance the invasion of new tissues. (Liotta L. A., Scientific American Feb: 54 (1992); and Liotta L. A. and Stetler-Stevenson W.G., Cancer Biol. 1: 99 (1990)). Most deaths incurred from cancer are due to its metastatic spread to secondary organs, therefore an inhibitor of metastasis would have widespread therapeutic use.

Cathepsin D is also believed to be associated with degenerative brain changes, such as those associated with Alzheimer's disease. Cathepsin D is associated with cleavage of amyloid- β -protein precursor, (Cataldo, A.M. et al., Proc. Natl. Acad. Sci. 87: 3861(1990)) including a mutant precursor that enhances amyloid protein production in transfected cells (Ladror, U.S., et al. J. Biol. Chem. 269: 18422

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(1994)). The amyloid- β -protein that results from proteolysis of the amyloid- β -protein precursor leads to senile plaque formation in brains and may be responsible for Alzheimer's disease. Recently elevated levels of cathepsin D have been found in cerebrospinal fluid in Alzheimer's patients (Schwager, A.L., et al. J. Neurochem. 64: 443 (1995)).

There is little known about substrate specificity and specific inhibitors for cathepsin D (Agarwal, N.S. and Rich, D.H., J. Med. Chem. 29: 2519 (1986); Jupp, R.A. et al., Biochem. J. 265: 871 (1990); Scarborough, P.E. et al., Protein Science 2: 264 (1993); Baldwin, E.T. et al., Proc. Natl. Acad. Sci. 90: 6796 (1993)).

Accordingly, there is a need for the synthesis of compounds which are specific for the inhibition of the activity of cathepsin D and which may be used in the treatment of metastatic disease and Alzheimer's disease.

A number of cathepsin D inhibitors have been reported (Lin, T.Y. and Williams, H.R., J. Biol. Chem. 25: 11875 (1970)). Agarwal and Rich reported the design and synthesis of cathepsin D inhibitors wherein the scissile dipeptide unit in a substrate sequence was replaced with a statine (Sta) residue or by a phenylstatine (Pst) unit. (Agarwal, N.S. and Rich, D.H., J. Med. Chem. 29: 2519 (1986)). Further, Agarwal and Rich evaluated the inhibition of cathepsin D by various analogues of pepstatin, finding that the fragment spanning P₄ to P', is necessary for the maximum inhibition of bovine cathepsin D.

U.S. Patent No. 4,746,648 describes peptide derivatives, modeled on the basis of pepstatin, which inhibit renin and acid protease. Tamburini et al., EP 0 569 777 A2 relates to the use of cathepsin D inhibitors for Alzheimer's disease. Patents relating to cyclic peptide inhibitors of renin include Boger et al., U.S. Patent No. 4,489,099 and Watkins, U.S. Patent No. 4,906,613.

Cyclizing peptidomimetics can increase binding to a target enzyme due to preorganization into the desired conformation. Additionally, such macrocycles offer increased stability against proteolytic cleavage. An area where this approach has been extensively explored is that of renin inhibitors, where various cycles connecting

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different positions were introduced. The most successful approach was cyclizing from the P₂ to P'₁ position to generate a series of potent, orally bioavailable renin inhibitors (Weber, A.E. et al. J. Med. Chem. 34: 2692 (1991); Dhanoa, D.S. et al. Tet. Lett. 33: 1725 (1992); Weber, A.E. et al. J. Med. Chem. 35: 3755 (1992); Yang, L. et al. Tet. Lett. 34: 7035 (1993)). Cyclization of P₄ to P₂ also gave potent renin inhibitors (Sham, H.L. et al. J. Chem. Soc. Chem. Commun. 666 (1990); Thaisrivongs, S. et al. J. Med. Chem. 34: 1276 (1991)). Some other cycles have been studied (Szewczuk, Z. et al. Int. J. Pept. Prot. Res. 40: 233 (1992); Sham, H.L. et al. J. Med. Chem. 31: 284 (1988); Dutta, A.S. et al. J. Med. Chem. 33: 2552 and 2560 (1990)), but the only example of which we are aware of a P₂ to P'₃ bridge in an aspartyl protease inhibitor is a simple disulfide (Boger, J. Peptides 1983, pp. 569-578, Proceedings of the 8th American Peptide Symposium).

The cyclic compounds described herein differ significantly from this prior art. Many of the compounds incorporate a P₂ to P'₃ cycle where the scissile bond isostere is part of the ring. Since the macrocycle spans a large portion of the binding cleft, activity is retained after truncation to remove the exocyclic backbone extension. This has the dual advantages of decreasing molecular weight and removing the residues most subject to metabolic cleavage.

Modeling studies reveal that the active sites of the aspartic hemoglobins (Plasmeppsins I and II) from *Plasmodium falciparum* are highly homologous with that of human cathepsin D. Goldberg, et al. isolated and characterized plasmeppsins I and II, aspartic proteases responsible for the initial cleavage of hemoglobin that occurs inside the protozoan *Plasmodium*'s digestive vacuole (Goldberg, D.E. et al., J. Exp. Med. 173: 961 (1991), Hill, J. et al. FEBS Letters 352: 155 (1994)). Protozoans of the genus *Plasmodium* are the causative agents of malaria. The cleavage of hemoglobin by plasmeppsins I and II occurs at sites within the hemoglobin sequence that are conserved in human hemoglobins. These cleavage events are essential for the conformational breakdown of hemoglobin that enables its subsequent cleavage by a series of other proteolytic enzymes. The digested hemoglobin is a primary nutrition source for the malarial parasite, which cannot grow

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in the absence of functional hemoglobins. Goldberg has demonstrated that inhibitors of plasmepsins can kill the parasite in a cell culture of infected human erythrocytes (Goldberg, D.E. et al., EMBO J. 13: 306 (1994), Gluzman, I.Y. et al. J. Clin. Invest. 93: 1602 (1994))

Therefore, there is a need for cathepsin D inhibitors in the treatment of Alzheimer's disease, cancer, and for aiding in the further elucidation of the roles of cathepsin D in human diseases and a need for plasmepsin inhibitors to treat malaria.

It is an object of the present invention to provide cathepsin D and plasmepsin I and II inhibitors for use in the treatment of metastatic disease, for use in the inhibition of cleavage of β -amyloid precursors and for the prevention of progressive neurological dysfunction in Alzheimer's disease, as well as for the prevention of the growth of *Plasmodium* parasites including *Plasmodium falciparum*, the most deadly cause of malaria.

SUMMARY OF THE INVENTION

The present invention relates to cathepsin D and plasmepsin I and II inhibitor compounds and their use as pharmaceutically active agents. The present invention further provides for uses of these inhibitors for the prevention and treatment of diseases, such as, for example, cancer, Alzheimer's disease, and malaria.

Specifically, the present invention provides for novel linear compounds as well as branched and cyclic analogs of these compounds which are inhibitors of cathepsin D and which exhibit inhibitory potency against plasmepsins I and II. Further, the present invention relates to novel related cyclic compounds which are cathepsin D and plasmepsin I and II inhibitors.

The present invention also provides for pharmaceutical compositions comprising effective amounts of at least one of the present inhibitors for use in the prevention and treatment of diseases, such as cancer, Alzheimer's disease, and malaria.

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Specifically, the present invention provides for pharmaceutical compositions which decrease the levels of activity of cathepsin D present in a subject, thereby inhibiting cancer cell invasion and metastasis.

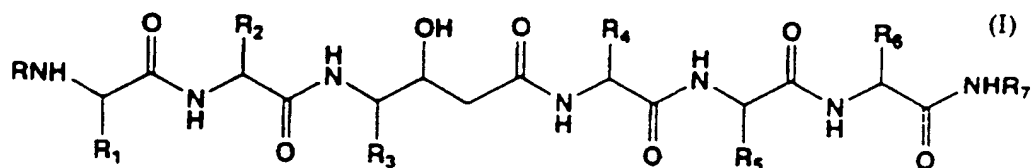
The present invention also provides for pharmaceutical compositions comprising at least one of the present inhibitors for the treatment and prevention of Alzheimer's disease which decrease the occurrence of cleavage of amyloid- β -protein precursors and senile plaque formation in a subject.

Further, the present invention provides for pharmaceutical compositions which inhibit plasmepsin I or II or both and prevent hemoglobin degradation and are thus useful in the treatment of malaria.

These and other features of the invention will be better understood through the following detailed description of the invention. The scope of the invention is limited only through the claims appended hereto.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to compounds of formula (I):



in which R, R₇ = represent hydrogen, alkoxycarbonyl, aralkoxycarbonyl, alkylcarbonyl, cycloalkylcarbonyl, cycloalkylalkoxycarbonyl, cycloalkylalkanoyl, aralkanoyl, aroyl, aryloxycarbonyl, aryloxycarbonylalkyl, aryloxyalkanoyl, heterocyclylcarbonyl, heterocyclyloxycarbonyl, heterocyclylalkanoyl, heterocyclylalkoxycarbonyl, heteroaralkanoyl, heteroaralkoxycarbonyl, heteroaryloxy-carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxyalkyl, heteroaryloxyalkyl, hydroxyalkyl, aminocarbonyl, aminoalkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted aminoalkyl, alkoxyalkyl.

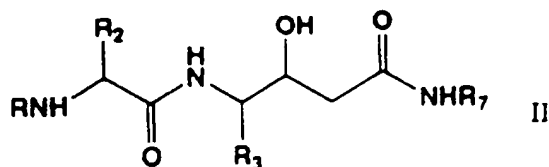
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alkylthioalkyl, mono- and disubstituted aminoalkanoyl radicals wherein the substituents are selected from alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkylalkyl radicals, or where said aminoalkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

R_1 , R_2 , R_4 , R_5 , R_6 = the side chain of a residue of an amino acid remaining after formation of a peptide linkage and include optionally substituted lower alkyl, lower cycloalkyl, aryl, aralkyl and heteroaryl;

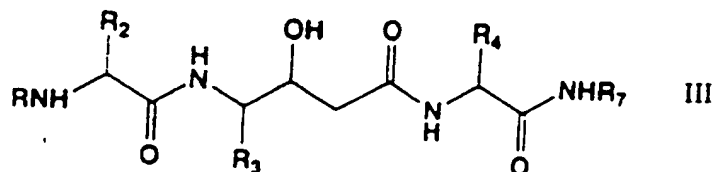
R_3 = optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, aralkyl, aralkoxy, aralkylthio, aralkylamino, cycloalkylalkyl, cycloalkylalkoxy, cycloalkylalkylthio, cycloalkylalkylamino. Also encompassed are compounds where R_2 and R_3 or R_1 and R_3 are connected by a bridging moiety having 4-14 atoms comprised of any stable combination of C, N, O, or S. This chain may be optionally substituted by halo, hydroxy, amino, lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, oxo, thiono, alkylimino, mono- or dialkylmethylidene, COR_3 . The bridging group may also be unsaturated so as to include the residues of alkenes, imines, alkynes and allenes. Furthermore, any part of the bridging moiety may comprise part of an optionally substituted aromatic, heteroaromatic, or cycloalkyl ring. Amino acids from which the residues containing R_1 , R_2 , R_4 , R_5 and R_6 can be derived include glycine, alanine, valine, leucine, isoleucine, serine, threonine, cysteine, methionine, aspartic acid, asparagine, glutamic acid, glutamine, lysine, hydroxylysine, arginine, phenylalanine, tyrosine, tryptophan, and histidine. Useful substituents and the optional substituted R_1 , R_2 , R_4 , R_5 and R_6 moieties include the functional groups which are attached to the above named aminoacids.

The present invention further relates to compounds of the Formula (II):



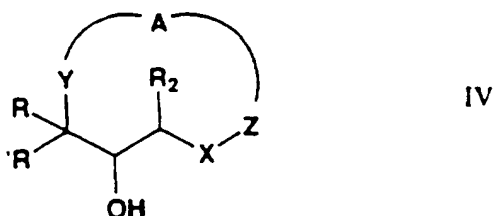
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wherein R, R₂, R₃, R₄, R₇ are defined as in formula I, or of Formula (III):



in which R, R₂, R₃, R₄, R₇ are defined as in formula I.

The present invention also relates to cyclic compounds of formula (IV):



where X = CWNR, CWOR, S(O)_nNR, P(O)(Q)NR; W = O, S, NR; n = 0, 1, 2;

Q = R, OR, N(R)₂

Y = NRCW, NRS(O)_n, NRP(O)(Q)

Z = CRR', NR

A = a bridging group having 2-15 atoms comprised of any stable combination of C, N, O, or S. This bridging group may itself be bridged by one or more chains comprised of C, N, O, or S atoms so as to generate additional rings of 3-7 atoms. The bridging groups may be optionally substituted by OH, NH₂, halo, optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, oxo, thiono, alkylimino, mono- or dialkylmethylidene, COR₂. The bridging moiety may also be unsaturated so as to contain portions which are the residues of alkene, imine, alkyne and allene groups. Furthermore, any part of the bridging moiety may comprise part of an optionally substituted aromatic, heteroaromatic, alicyclic, or heterocyclic ring.

R = is as defined hereinabove

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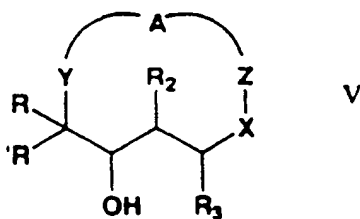
$R_1, R_2 = H$, halo, optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino

$R_3 =$ is as defined hereinabove

Also encompassed within the scope of the invention are compounds of formula IVa where IVa has the structure shown for IV and $X, Y = NRCW, NRS(O)_n, NRP(O)(Q)$; $Z = CRR', NR, O, S$ and all other definitions are as above.

Additionally contemplated are compounds of the formula IVb where IVb has the structure shown for IV, where $X = OCWNR, SCWNR$; $Y = NRCW, NRS(O)_n, NRP(O)(Q)$; $Z = CRR', NR$ and all other definitions are as above.

The present invention also relates to cyclic compounds of formula (V):



where $X = CWNR, S(O)_nNR, P(O)(Q)NR$; $W = O, S, NR$; $n = 0, 1, 2$;

$Q = R, OR, N(R)_2$

$Y = NRCW, NRS(O)_n, NRP(O)(Q)$

$Z = CRR', NR$

$A =$ is as defined hereinabove

$R =$ is as defined hereinabove.

R_1, R_2 and $R_3 = H$, halo, optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino;

Also encompassed within the present invention are compounds of formula Va where Va has the structure shown for V and where $X, Y = NRCW, NRS(O)_n, NRP(O)(Q)$; $Z = CRR', NR, O, S$ and all other definitions are as above.

Additionally encompassed are compounds of formula Vb where Vb has the structure shown for V and where $X = OCWNR, SCWNR$; $Y = NRCW, NRS(O)_n, NRP(O)(Q)$; $Z = CRR', NR$ and all other definitions are as above.

As utilized herein, the term "alkyl", alone or in combination, means a straight-chain or branched-chain alkyl radical containing from 1 to about 18, preferably from 1 to about 10, carbon atoms. Examples of such radicals include methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, amyl, isoamyl, hexyl, octyl and the like. The term "alkenyl", alone or in combination, means a straight-chain or branched-chain hydrocarbon radical having one or more double bonds and containing from 2 to about 18 carbon atoms preferably from 2 to about 10 carbon atoms. Examples of suitable alkenyl radicals include ethenyl, propenyl, allyl, 1,3-butadienyl and the like. The term "alkynyl", alone or in combination, means a straight-chain hydrocarbon or branched chain radical having one or more triple bonds and containing from 2 to about 10 carbon atoms. Examples of alkynyl radicals include ethynyl, propynyl, propargyl, butynyl and the like. The term "alkoxy", alone or in combination, means an alkyl ether radical wherein the term alkyl is as defined above. Examples of suitable alkyl ether radicals include methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, iso-butoxy, sec-butoxy, tert-butoxy and the like. The term "cycloalkyl", alone or in combination, means a saturated or partially saturated monocyclic, bicyclic or tricyclic alkyl radical wherein each cyclic moiety contains from about 3 to about 8 carbon atoms. Examples of such cycloalkyl radicals include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and the like. The term "cycloalkylalkyl" means an alkyl radical as defined above which is substituted by a cycloalkyl radical as defined above. Examples of cycloalkylalkyl radicals include cyclopropylmethyl, cyclopropylethyl, cyclohexylmethyl, cyclohexylethyl and the like. The term "aryl", alone or in combination, means an aromatic monocycle or bicyclic or tricyclic such as phenyl, naphthyl, or anthracenyl which optionally carries one or more substituents selected from alkyl, alkoxy, halogen, hydroxy, amino, nitro, cyano, haloalkyl and the like, such as phenyl, p-tolyl, 4-ethoxyphenyl, 4-(tert-butoxy) phenyl, 4-fluorophenyl, 4-chlorophenyl, 4-hydroxyphenyl, 1-naphthyl, 2-naphthyl, and the like. The term "aralkyl", alone or in combination, means an alkyl radical as defined above in which one hydrogen atom is replaced by an aryl radical as defined above, such as benzyl, 2-phenylethyl and the like. The term "aryloxy" means a radical of the

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formula aryl-O- in which the term aryl has the significance given above. The term "alkanoyl", alone or in combination, means an acyl radical derived from an alkanecarboxylic acid, examples of which include acetyl, propionyl, butyryl, valeryl, 4-methylvaleryl, and the like. The heterocyclyl portion of a heterocyclyl-containing group is a saturated or partially unsaturated monocyclic, bicyclic or tricyclic heterocycle which contains one or more hetero atoms selected from nitrogen, oxygen and sulphur, which is optionally substituted on one or more carbon atoms by halogen, alkyl, alkoxy, oxo, and the like, and/or on a secondary nitrogen atom by alkyl, aralkoxycarbonyl, alkanoyl, aryl or aralkyl or on a tertiary nitrogen atom by oxido and which is attached via a carbon atom. The heteroaryl group is an aromatic monocyclic, bicyclic, or tricyclic heterocycle which contains the hetero atoms and is optionally substituted as defined above with respect to the definition of heterocyclyl. Examples of such heterocyclyl and heteroaryl groups are pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl, thiomorpholinyl, pyrrolyl, imidazolyl (e.g. imidazol-4-yl, 1-benzoyloxycarbonylimidazol-4-yl, etc.), pyrazolyl, pyridyl, pyrazinyl, pyrimidinyl, furyl, thienyl, triazolyl, oxazolyl, thiazolyl, indolyl (e.g. 2-indolyl), quinolinyl, (e.g., 3-quinolinyl, 2-quinolinyl, etc.), isoquinolinyl (e.g., 1,2,3,4-tetrahydro-1-oxo-isoquinolinyl, etc.), quinoxalinyl, β -carbolinyl, benzofuranyl, 1-, 2-, 4- or 5-benzimidazolyl, and the like. The term "aminoalkanoyl" means an acyl group derived from an amino-substituted alkanecarboxylic acid wherein the amino group can be a primary, secondary or tertiary amino group containing substituents selected from hydrogen, alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl radicals and the like. The term "halogen" means fluorine, chlorine, bromine or iodine. The term "haloalkyl" means an alkyl radical having the significance as defined above wherein one or more hydrogens are replaced with a halogen. Examples of such haloalkyl radicals include chloromethyl, 1-bromoethyl, fluoromethyl, difluoromethyl, trifluoromethyl, 1,1,1-trifluoroethyl and the like.

The possible optional substituents mentioned in the hereinabove generic description include at least one alkyl, cycloalkyl, aryl, aralkyl, alkaryl, heteroaryl, alkoxy, halogen, hydroxy, amino, nitro, cyano, haloalkyl wherein the optional

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substituents may also be optionally substituted and the radicals which are optionally substituted may be singly or multiply substituted with the same or different optional substituents.

The compounds provided by the present invention are advantageous as demonstrated by their activity. Furthermore, the structures of the cyclic compounds of the present invention potentially provide protection of the compounds from enzyme degradation. The low molecular weight and few peptide bonds in these analogs may also contribute to improved bioavailability.

Specific, but non-limiting examples of peptides of Formula (I) useful in the present invention include the following:

		K_i cathepsin D (nM)
(1)	Iva-Val-Val-Sta-Val-Leu-Gly-NH ₂ (SEQ. ID. NO. 1);	0.1
(2)	Iva-Gln-Val-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 2);	0.35
(3)	Iva-Lys-Val-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 3);	3.9
(4)	Tba-Val-Val-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 4);	0.04
(5)	Iva-Val-Ile-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 5);	0.04
(6)	Iva-Val-Leu-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 6);	0.5
(7)	Tba-Val-Val-Pst-Val-Leu-Gly-NH ₂ (SEQ. ID. NO. 7);	0.015
(8)	Tba-Val-Cys-Pst-Val-Leu-Gly-NH ₂ (SEQ. ID. NO. 8);	0.25
(9)	Tba-Val-Glu-Pst-Val-Leu-Gly-NH ₂ (SEQ. ID. NO. 9);	1.25
(10)	Tba-Val-Asp-Pst-Val-Leu-Gly-NH ₂ (SEQ. ID. NO. 10);	2.1
(11)	Iva-Val-Val-Sta-Ala-Leu-Gly-NH ₂ (SEQ. ID. NO. 11);	0.03
(12)	Tba-Val-Cys-Pst-Val-Cys-Gly-NH ₂ (SEQ. ID. NO. 12);	0.66

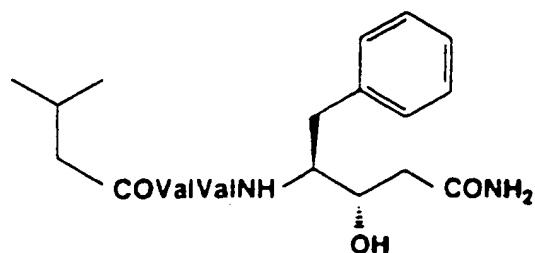
which has a PMI K_i = 32 nM PMII K_i = 1.7 nM.

Iva = isovaleryl

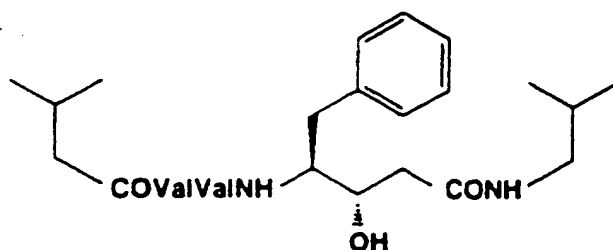
Tba = t-butylacetyl

Specific, but not limiting examples of peptides of Formula (II) of the present invention include the following:

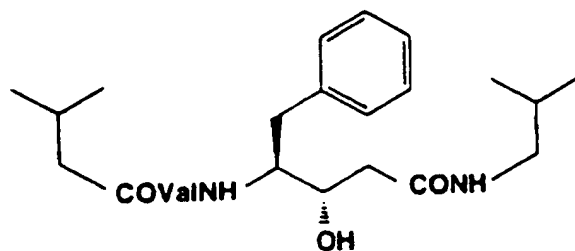
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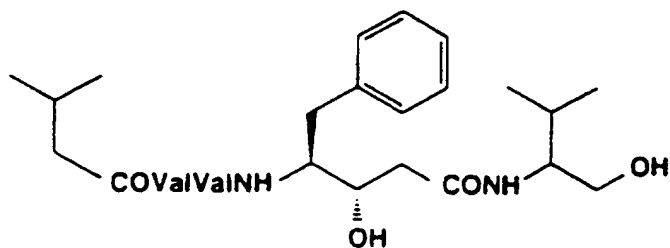
which has a K_i of 800 nM, (SEQ. ID. NO. 13);



which has a K_i of 12 nM, (SEQ. ID. NO. 14);

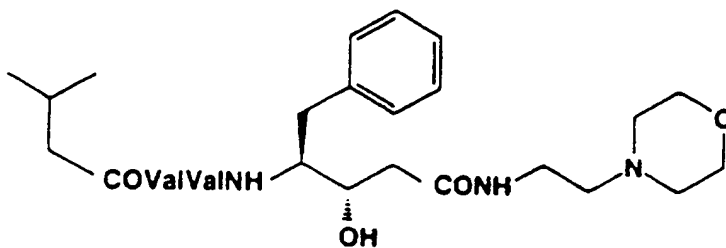


which has a K_i of 1600 nM, (SEQ. ID. NO. 15);



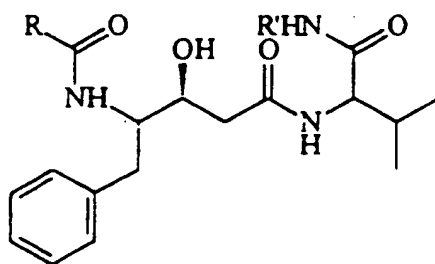
which has an IC_{50} of 600 nM. (SEQ. ID. NO. 16);

- 13 -



which has a K_i of 1830 nM (SEQ.ID.NO. 17);

Additional Examples of Cathepsin D Inhibitors within Formula II are shown below based on the formula:

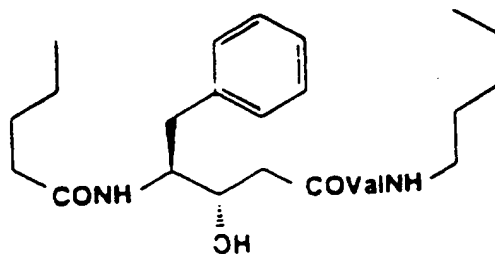


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R	R ¹	MW	K _i (nM)
(CH ₂) ₄ CH ₃	(CH ₂) ₆ CH ₃	503	24
(CH ₂) ₂ SPr	(CH ₂) ₇ SPr	539	75
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₂ SCH ₂ Ph	635	5
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₃ SCH ₂ Ph	649	17
(CH ₂) ₃ SCH ₂ Ph	(CH ₂) ₃ SCH ₂ Ph	663	4.0
O(CH ₂) ₃ -4-Pyr	(CH ₂) ₃ SCH ₂ Ph	634	249
(CH ₂) ₃ SCH ₂ Ph	(CH ₂) ₆ CH ₃	597	6.2
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₆ CH ₃	583	7.7
(CH ₂) ₄ CH ₃	(CH ₂) ₃ SBn	569	6.1
(CH ₂) ₂ CO ₂ H	n-C ₇ H ₁₅	505	41
(CH ₂) ₂ CO ₂ Me	n-C ₇ H ₁₅	519	178
(CH ₂) ₂ CONH ₂	n-C ₇ H ₁₅	504	479
(CH ₂) ₃ CO ₂ H	n-C ₇ H ₁₅	519	22
(CH ₂) ₃ CO ₂ Me	n-C ₇ H ₁₅	533	112
CH ₂ -p-Ph-Ph	n-C ₇ H ₁₅	599	16
(E)-3-CH=CHbenzodioxolane	n-C ₇ H ₁₅	579	45

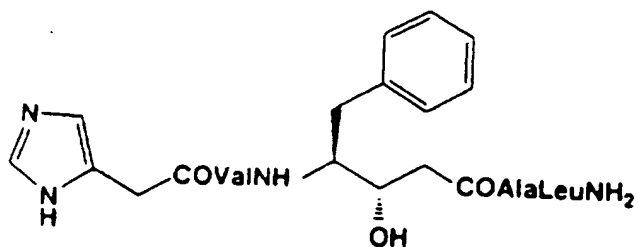
Ph = phenyl, pyr = pyridyl and Bn = benzyl

Specific, but not limiting examples of compounds of Formula (III) include:

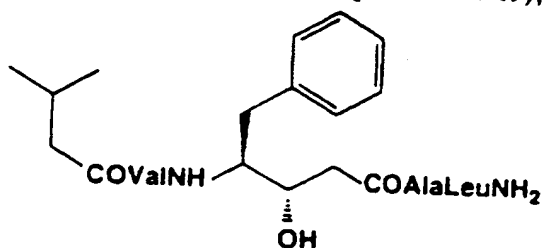


which has an K_i of 400 nM, (SEQ. ID. NO. 18);

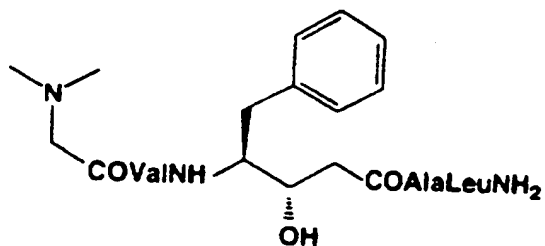
- 15 -



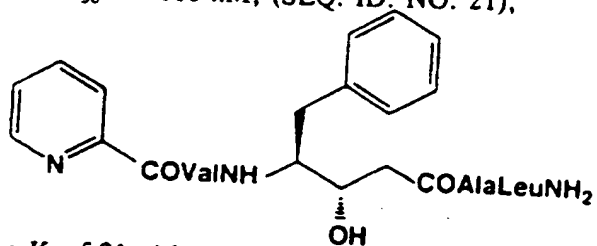
which has an IC₅₀ of 500 nM, (SEQ. ID. NO. 19);



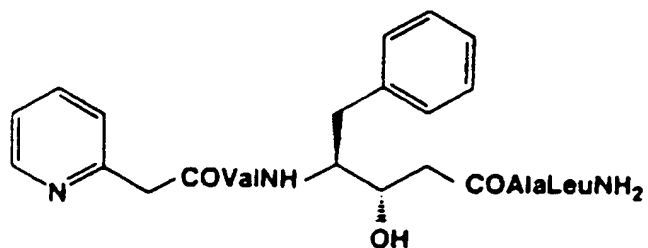
which has a K_i of 8 nM, (SEQ. ID. NO. 20);



which has an IC₅₀ of 1000 nM, (SEQ. ID. NO. 21);

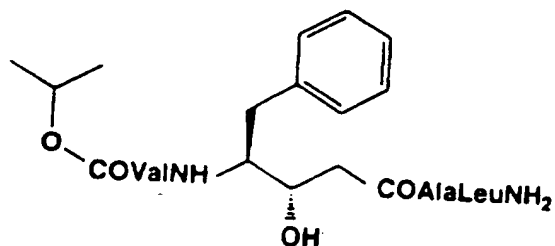


which has a K_i of 21 nM, (SEQ. ID. NO. 22);

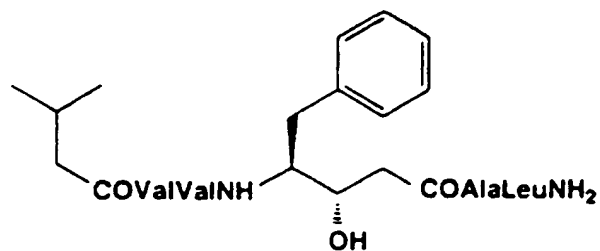


which has a K_i of 20 nM, (SEQ. ID. NO. 23);

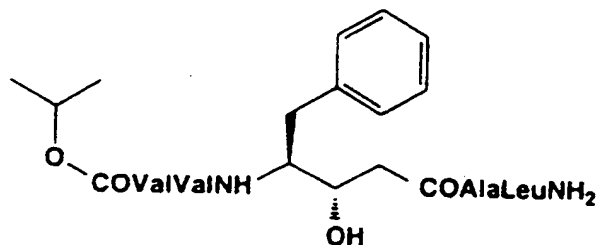
- 16 -



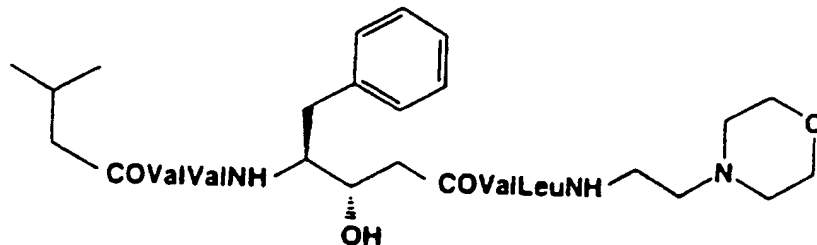
which has an IC_{50} of 20 nM, (SEQ. ID. NO. 24);



which has a K_i of 0.18 nM, (SEQ. ID. NO. 25);



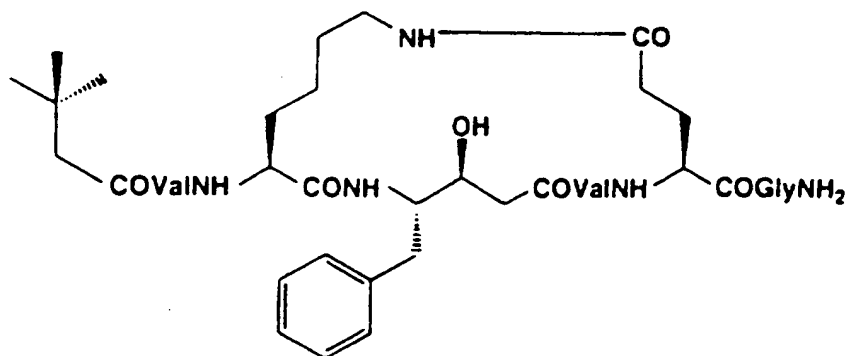
which has a K_i of 0.24 nM, (SEQ. ID. NO. 26);



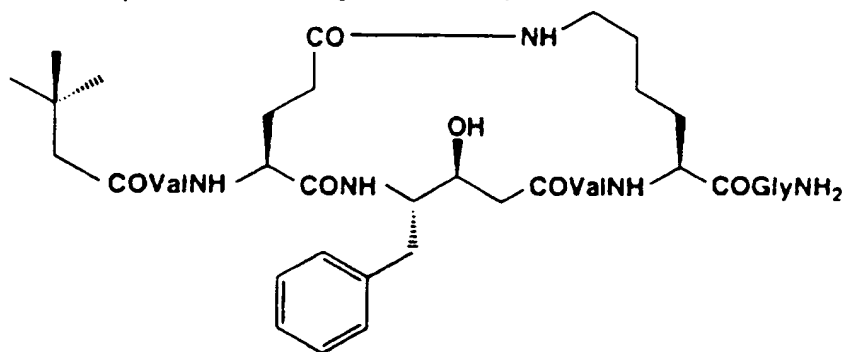
which has a K_i of 0.14 nM, (SEQ. ID. NO. 27).

Specific examples of cyclic compounds within the scope of the present invention include: (Formula I:)

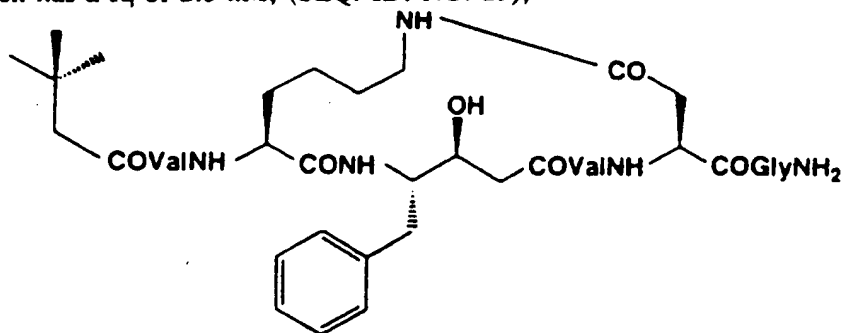
- 17 -



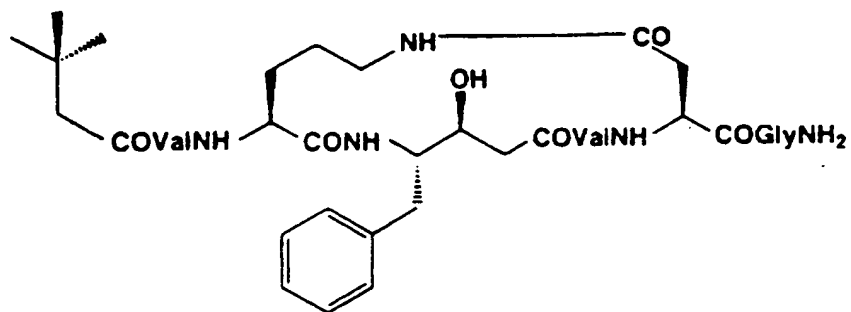
which has a K_i of 1.4 nM, (SEQ. ID. NO. 28);



which has a K_i of 2.3 nM, (SEQ. ID. NO. 29);

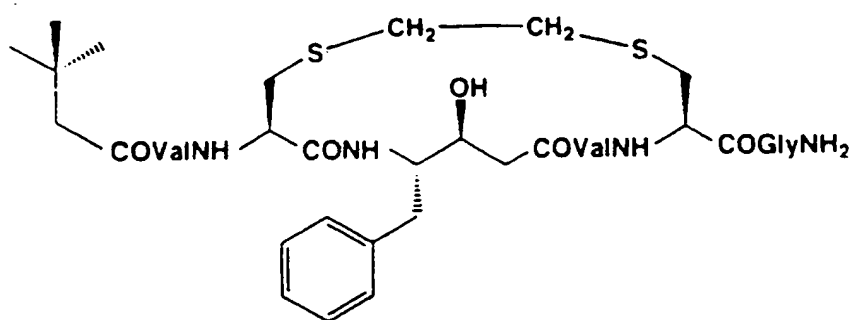


which has a K_i of 12 nM, (SEQ. ID. NO. 30);

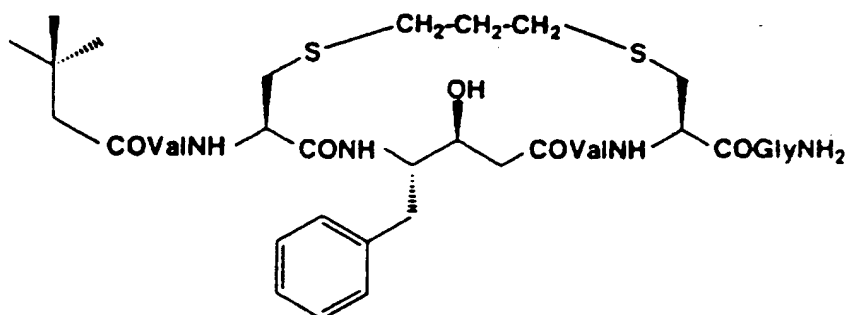


which has a K_i value of 100 nM, (SEQ. ID. NO. 31);

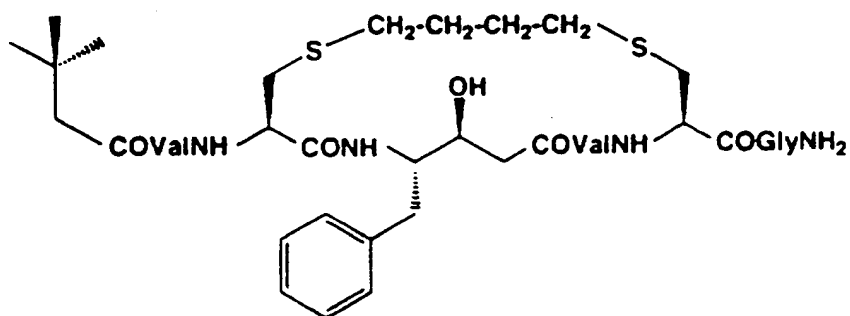
- 18 -



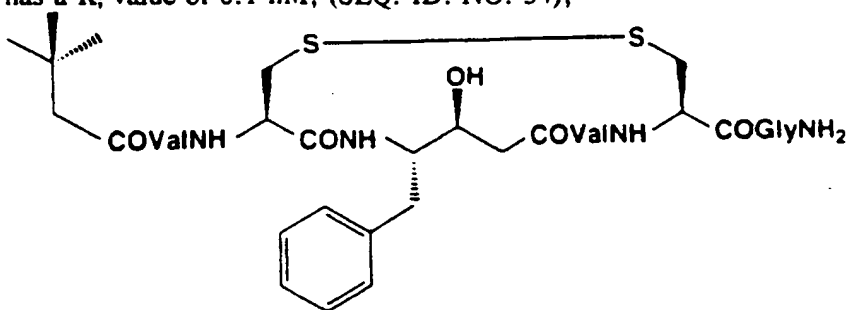
which has a K_i value of 1.9 nM, (SEQ. ID. NO. 32);



which has a K_i value of 1.5 nM, (SEQ. ID. NO. 33);

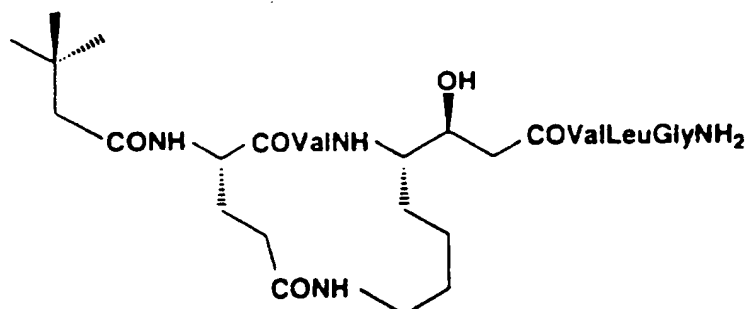


which has a K_i value of 0.1 nM, (SEQ. ID. NO. 34);

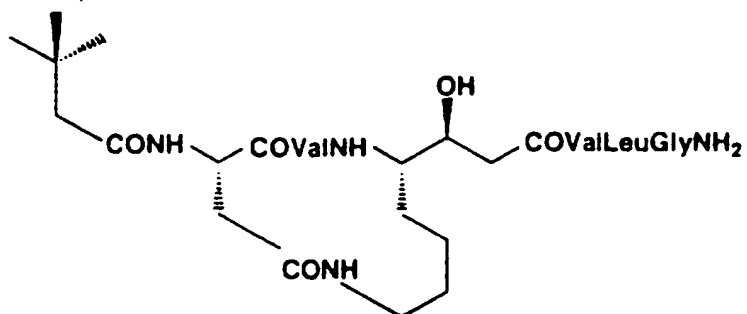


which has a K_i value of 150 nM, (SEQ. ID. NO. 35);

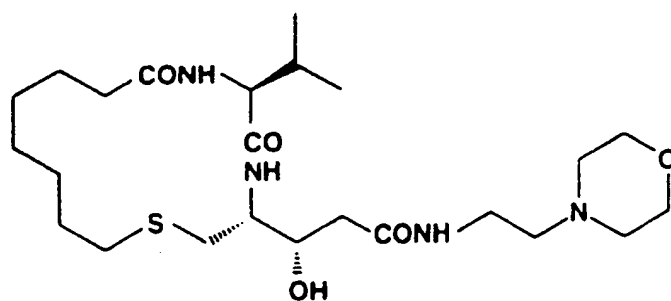
- 19 -



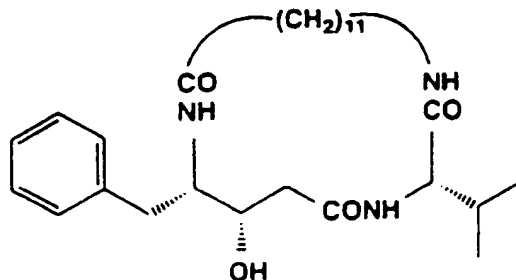
which has a K_i value of 0.26 nM, (SEQ. ID. NO. 36);



which has a K_i value of 2.5 nM, (SEQ. ID. NO. 37);
Formula II:

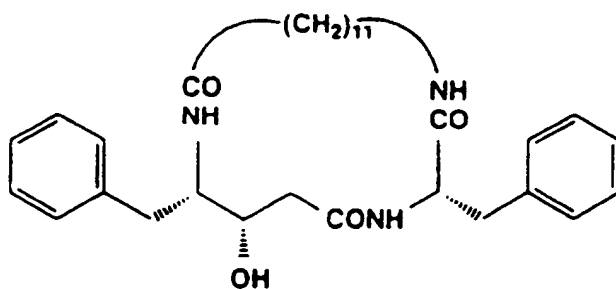


which has an IC_{50} of 10,000 nM (SEQ.ID.NO. 38);
Formula IV:

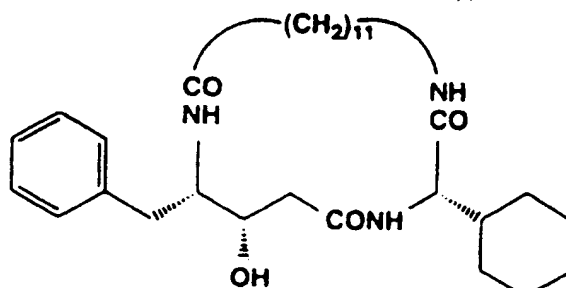


which has a K_i value of 4 nM; (SEQ.ID.NO. 39);

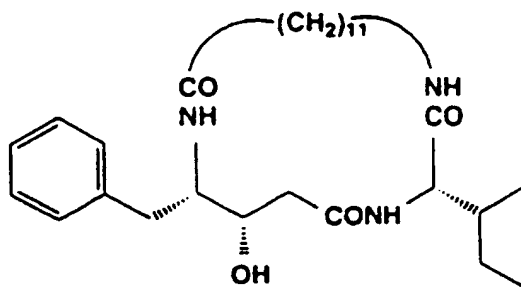
- 20 -



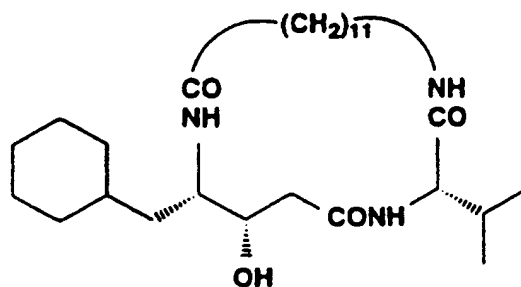
which has a K_i value of 21 nM; (SEQ.ID.NO. 40);



which has a K_i value of 85 nM; (SEQ.ID.NO. 41);

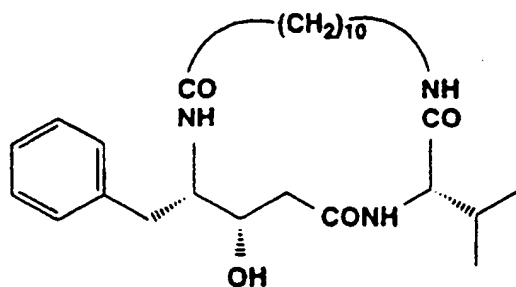


which has a K_i value of 22 nM; (SEQ.ID.NO. 42);

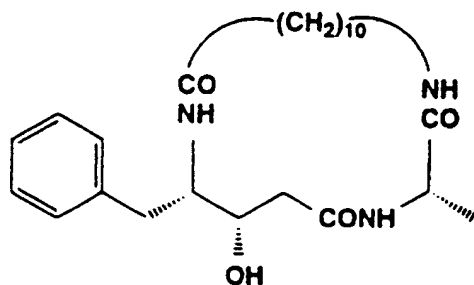


which has a K_i value of 10 nM; (SEQ.ID.NO. 43);

- 21 -

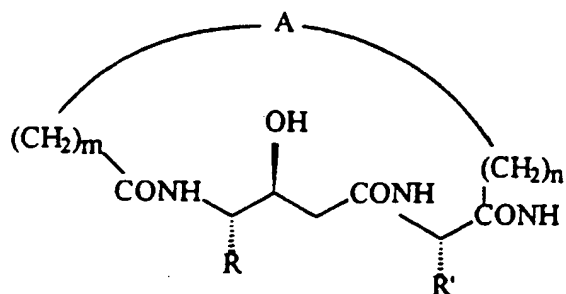


which has a K_i value of 20 nM; (SEQ.ID.NO. 44);



which has a K_i value of 140 nM. (SEQ.ID.NO. 45).

Additional Examples of Cathepsin D Inhibitors within Formula IV are shown below based on the following formula:



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R	R ¹	m	n	A	Ki [nM]
PhCH ₂	D-iPr	11	-	-	-
PhCH ₂	iBu	11	-	-	20
PhCH ₂	Me	11	-	-	46
PhCH ₂	Et	11	-	-	22
PhCH ₂	3-PyrCH ₂	11	-	-	38
PhCH ₂	CH ₃ CH(OCH ₂ Ph) CH ₂	11	-	-	460
PhCH ₂	CH ₃ CH(OH)CH ₂	11	-	-	300
PhCH ₂	tBu	11	-	-	18
PhCH ₂	Ph	11	-	-	11
p-(PhCH ₂ O)- PhCH ₂	iPr	11	-	-	41
p-(OH)PhCH ₂	iPr	11	-	-	53, 45, 27
p- (OMe)PhCH ₂	iPr	11	-	-	8.3
p-(OEt)- PhCH ₂	iPr	11	-	-	8.1
p-(OiPr)- PhCH ₂	iPr	11	-	-	7.4
p-(3PyrCH ₂ O)- PhCH ₂	iPr	11	-	-	81
p- MeO(CH ₂) ₂ O- PhCH ₂	iPr	11	-	-	13
(F ₃ -Ph)CH ₂	iPr	11	-	-	90
(m,p- Cl ₂ Ph)CH ₂	iPr	11	-	-	-
(mp-F ₂ Ph)CH ₂	iPr	11	-	-	-
p-F-PhCH ₂	iPr	11	-	-	-
HSCH ₂	iPr	11	-	-	410

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R	R ¹	m	n	A	K _i [nM]
MeSCH ₂	iPr	11	-	-	150, 183
EtSCH ₂	iPr	11	-	-	84
PrSCH ₂	iPr	11	-	-	90
CH ₂ =CHCH ₂ SCH ₂	iPr	11	-	-	140
BuSCH ₂	iPr	11	-	-	-
iBuSCH ₂	iPr	11	-	-	-
PhCH ₂ SCH ₂	iPr	11	-	-	68
PhCH ₂ CH ₂	iPr	11	-	-	1600
2-naph-CH ₂	iPr	11	-	-	1.1
1-naph-CH ₂	iPr	11	-	-	121
4-thiazolyl- CH ₂	iPr	11	-	-	186
3-indolyl-CH ₂	iPr	11	-	-	45
PhCH ₂	iPr	-	10	O	36
PhCH ₂	iPr	-	12	O	10
PhCH ₂	iPr	2	2	S(CH ₂) ₃ S	1900
PhCH ₂	iPr	2	2	S(CH ₂) ₄ S	573
PhCH ₂	iPr	2	2	S(CH ₂) ₅ S	53
PhCH ₂	iPr	2	2	S(CH ₂) ₆ S	19
PhCH ₂	iPr	2	2	S(CH ₂) ₇ S	56
PhCH ₂	iPr	2	2	SCH ₂ CH=CHCH ₂ S (cis)	380
PhCH ₂	iPr	2	2	SCH ₂ CH=CHCH ₂ S (trans)	320
PhCH ₂	iPr	2	2	SCH ₂ CCCH ₂ S	65
PhCH ₂	iPr	2	2	SCH ₂ p-Ph-CH ₂ S	9.8
PhCH ₂	iPr	2	2	SCH ₂ m-Ph-CH ₂ S	
PhCH ₂	iPr	2	2	SCH ₂ o-Ph-CH ₂ S	

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R	R ¹	m	n	A	Ki [nM]
PhCH ₂	iPr	2	2	SCH ₂ P-(Cl) ₄ Ph- CH ₂ S	0.89
PhCH ₂	iPr	11	-	replace amide NH by O	63

naph. = naphthyl

The compounds of the present invention may exist in a free, i.e. unprotected, or protected form. The protected form herein refers to compounds wherein one or more reactive groups, e.g. amino groups or -OH groups, are substituted by a protecting group. Suitable protecting groups are any of those known in the art, such as acetyl, benzyloxycarbonyl and t-butoxycarbonyl.

The compounds of the present invention, whether they are in a free or protected form, may exist as salts or as complexes. Acid addition salts may be formed with organic acids, polymeric acids, and inorganic acids, for example. Such acid addition salt forms include *inter alia* the hydrochlorides and acetates. Complexes are herein defined as compounds of known type, formed on addition of inorganic substances, such as inorganic salts or hydroxides such as Ca- and Zn- salts, and/or on addition of polymeric organic substances.

The present invention further provides methods and compositions for preventing or treating diseases. Particular non-limiting examples of diseases include cancer including for example breast cancer, Alzheimer's disease, and malaria.

Specifically, this invention provides for the use of the compounds and compositions of the present invention to inhibit the activities of cathepsin D and plasmepsins I and II for treating and preventing diseases such as cancer, Alzheimer's disease, and malaria. The present invention also provides pharmaceutical compositions comprising the same.

The present invention further provides methods of preventing or treating a disease by the administration of a therapeutically effective amount of a cathepsin D or plasmepsin I or II inhibitor compound.

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More particularly, the present invention provides methods of treating cancer by administration of a therapeutically effective amount of at least one cathepsin D inhibitor described herein which, for example, inhibits the invasion and metastasis of cancerous cells.

In addition, the present invention provides methods of treating Alzheimer's disease by administration of a therapeutically effective amount of at least one cathepsin D inhibitor described herein which, for example, inhibits the formation of senile plaques.

In addition, the present invention provides methods of treating malaria by administration of a therapeutically effective amount of a plasmepsin I or II inhibitor described herein which, for example, inhibits the degradation of hemoglobin by the malarial intracellular parasite.

The present invention also provides methods of preventing or treating diseases by the administration of a therapeutically effective amount of at least one compound of the present invention in combination with chemotherapeutic agents, toxins, or irradiation. Examples of chemotherapeutic agents are known to those skilled in the art and include, but are not limited to, bleomycin, mitomycin, cyclophosphamide, doxorubicin, paclitaxel, and cisplatin.

In one embodiment of the invention, the compounds of the present invention are administered in a pharmaceutically acceptable carrier. A pharmaceutically acceptable carrier encompasses any of the standard pharmaceutical carriers such as sterile solution, tablets, coated tablets and capsules. Such carriers may typically contain excipients such as starch, milk, sugar, certain types of clay, gelatin, stearic acid, talc, vegetable fats or oils, gums, glycols, or other known excipients. Such carriers may also include flavor and color additives and other ingredients.

In the practice of the methods of this invention, the amount of the enzyme inhibitor incorporated in the composition may vary widely. Methods for determining the precise amount depend upon the subject being treated, the specific pharmaceutical carrier, the route of administration being employed, the frequency with

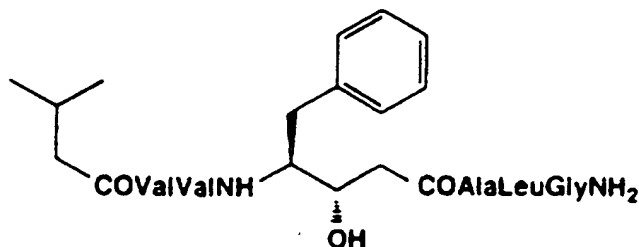
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which the compound is to be administered, and whether the composition is administered in conjunction with a chemotherapeutic agent and/or irradiation treatment.

The present invention further provides a method of treating a subject afflicted with a tumor which comprises contacting the tumor with an amount of one of the cathepsin D inhibitors described herein which is administered to the subject previous to, simultaneous to, or subsequent to, administration of a chemotherapeutic agent or to an amount of irradiation effective to treat the tumor. The administration of the composition may be effected by any of the well known methods, including but not limited to, oral, intravenous, intramuscular, and subcutaneous administration.

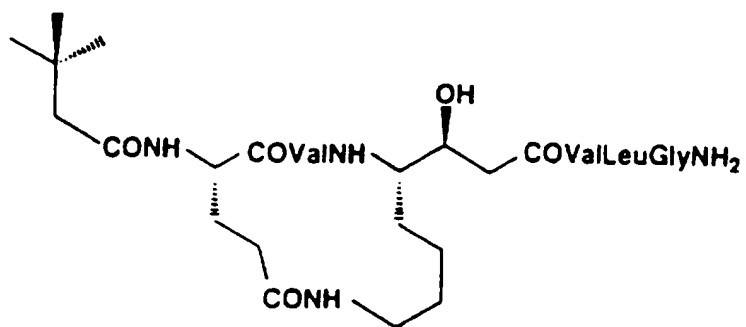
Plasmeprin I and II are enzymes which are required for specific degradation of hemoglobin by the malarial intracellular parasite. Due to the high active site similarity between cathepsin D and plasmeprins, some of the cathepsin D inhibitors of the present invention are useful in preventing the growth of *Plasmodium falciparum*, a causative agent of malaria. Non-limiting examples of compounds which exhibit plasmeprin I or II (aspartic hemoglobinses from *Plasmodium falciparum*) inhibitory activity include the following:

(Formula I):

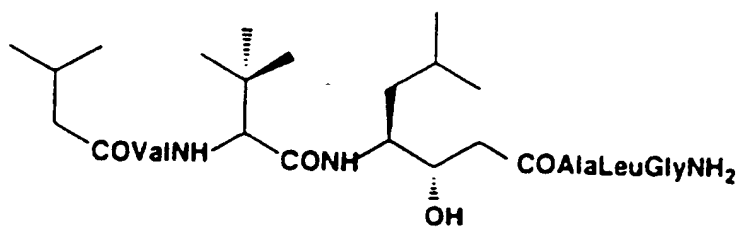


which has a PMI $K_i = 1.2$ nM PMII $K_i = 0.1$ nM (SEQ. ID. NO. 46);

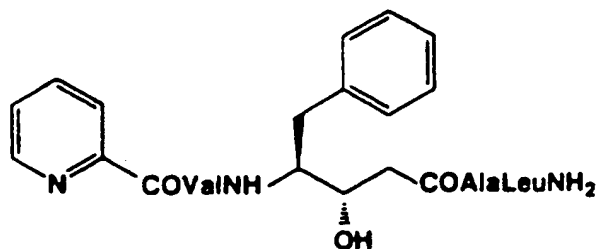
- 27 -



which has a PMI $K_i = 22$ nM PMII $K_i = 0.2$ nM (SEQ. ID. NO. 36);



which has a PMI $K_i = 7$ nM PMII $K_i = 3$ nM (SEQ. ID. NO. 47);
(Formula III):



which has a PMI $K_i = 16$ nM PMII $K_i = 0.9$ nm (SEQ.ID.NO. 22).

SYNTHESIS:**LINEAR INHIBITORS:**

The linear peptide inhibitors of the present invention were synthesized by the solid phase method using the Fmoc group for α -amino protection and acid labile groups for side chain protection of trifunctional amino acids e.g. [FmocCys(Trt)OH, FmocAsp(OtBu)OH and FmocGlu(OtBu)OH]. The aminomethyl polystyrene resin (Bachem California) was modified with acid labile linker Fmoc-2,4-dimethoxy-4'-(carboxymethoxy)-benzhydrylamine (Bachem Bioscience, Inc.) to a final substitution of 0.3-0.5 mmol/g. The Fmoc protected amino acids were coupled as HOBT esters, DIC (diisopropylcarbodiimide) was used for activation, and a 20% solution of piperidine in DMF (dimethyl formamide) for deprotection in each step. The Fmoc protected derivatives of statine and 3-hydroxy-4-amino-5-phenyl pentanoic acid (AHPPA or phenylstatine) were prepared according to the modified described procedure of Jouin, P. et al. J. Chem. Soc. Perkin. Trans. 1, 1177 (1987). The final cleavage of inhibitors from the resin was accomplished by treating with TFA (trifluoroacetic acid) containing 5% of water (and 3% of triethyl silane in the case of cysteine containing inhibitors). All the inhibitors were purified by reverse phase high performance liquid chromatography (HPLC) (column VYDAC C-18 2.5 X 25 cm) using water-acetonitrile mixtures containing 0.05% TFA in a gradient elution. Purity of all compounds was checked with analytical reverse phase HPLC (column CYDAC C-18 0.4 X 25 cm). All compounds gave correct molecular peaks in the mass spectrum (SIMS).

CYCLIC ANALOGS:**Preparation of Cyclic Disulfides.**

The dibenzylated dithiol (100-500 mg) was dissolved in 150 ml of refluxing anhydrous liquid ammonia. Sodium metal was then added in small pieces to the solution until it remained blue. The addition was finished when the blue color persisted for 3 minutes. The mixture was then decolorized with a crystal of NH_4Cl

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and ammonia was evaporated. The solid residue was suspended in 10% aq. KHSO_4 and the precipitated dithiol was filtered off. Yield is usually close to 100%.

The dithiol (30 mg) was dissolved in 30 ml of anhydrous degassed DMF containing 10 eq. of diisopropylethylamine (DIEA). A solution of 1.1 eq. of α,ω -dihaloalkane in 5 ml of DMF was then added dropwise during 12 hours to the above solution. The mixture was left to stand for an additional 12 hours at room temperature. DMF was evaporated under reduced pressure and the obtained solid residue was washed with 5% aq. NaHCO_3 and ether. Yield 70-90% of crude cyclic inhibitor. The product was purified by preparative HPLC for the purpose of K_i determination.

Other Cyclic Analogs

Compounds related to Seq. ID number 42 but lacking the Ile can be prepared by condensing an N-protected statine analog with an ω -amino ester, hydrolyzing the ester, removing the N-protecting group of the statine, and cyclizing to form the lactam. Alternatively an ester of statine can be condensed with an N-protected ω -amino acid to provide the N-acylated statine which can be deprotected and cyclized to give the lactam.

Analogs such as in compounds of formula IV where a carboxamide is replaced with a sulfonamide or a phosphonamide may be prepared in an analogous way to the carboxamides, but in place of typical peptide coupling conditions, more vigorous activation of the sulfonic or phosphoric acid such as by preparing the sulfonyl or phosphoryl chloride may be required.

The amides described above can be replaced by hydrazides by using a suitably C-protected ω -hydrazino acid and condensing it with a suitably activated carboxylic or related acid derivative.

The oxygen of the carbonyl containing derivatives such as carboxamides may be replaced by sulfur by using such common thionating agents as P_2S_5 or Lawesson's reagent. The resulting thiocarbonyls can be converted into imino

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derivatives by treating with amines, in some cases after prior activation of the thiocarbonyl by, for example, alkylation.

BRIDGING GROUPS

It is recognized that the ω -amino acid may have up to 4 of its carbon atoms replaced by heteroatoms such as O, N, or S. These amino acids may be prepared by using methods known to those skilled in the art. For example a shorter chain ω -halo acid derivative can be reacted with an N-protected amino alcohol, aminoalkanethiol, or diamine to provide an ω -amino acid with an O, S, or N beta to the amino terminus. Similarly an ω -hydroxy acid could be derivatized as, for example, a sulfonate, and a similar displacement reaction run. Alternatively a suitably protected shorter chain ω -amino, hydroxy, or sulfhydryl acid could be derivatized, for example, by reaction with a suitably N-protected amino alkyl halide or sulfonate. Similarly it is recognized that replacing a sulfonate or halide in the above alkylations by an epoxide would allow formation of chains with hydroxyl substituents. It should be noted that chains containing heteroatoms need not be synthesized only by using alkylation type chemistry. Chains containing an amide bond can be synthesized, for example, by reacting an amino acid with another amino acid using standard peptide coupling conditions to generate the ω -amino acid required. Furthermore an O-protected ω -amino acid could be reacted with a phosgene equivalent to provide an isocyanate which could then be reacted with an N-protected amino alcohol or diamine to provide a urethane or urea-substituted chain.

It is recognized that the chain need not be built up separately, but may be formed by attaching pieces of the chain to the amino and carboxy ends of the molecule and then connecting them in some way. This approach is exemplified in Seq. ID numbers 32-34, where the two cysteine substituents are connected by a dihalide to form the desired bridge compounds. Similarly Seq. ID. NO. 28 can be prepared by cyclization of a compound with an amino alkyl substituent at one end and a carboxy alkyl substituent at the other to give the chain containing an amide functionality.

It should be noted that chains where carbon atoms have been replaced by heteroatoms are amenable to further chemistry which can also lead to active compounds. For example sulfides can be converted by peracids to sulfoxides or sulfones. Secondary amines can be alkylated or acylated.

Chains containing oxo or hydroxy substitution can also be elaborated. For example oxo groups may be condensed with amines, hydrazines, hydroxylamine, alkoxyamines, phosphorous pentasulfide, diethylamino sulfur trifluoride (DAST), alkylidene triphenyl phosphoranes or the like to provide imines, hydrazones, oximes, thiones, difluoromethylenes, alkylidenes and the like. Reduction provides alcohols, or methylenes. Alcohols may be converted into halides for example by phosphorous trihalide, oxidized to oxo compounds, acylated to give esters or alkylated to give ethers.

The chain could also incorporate unsaturation such as one or more double or triple bonds. These can be generated by elaboration of saturated analogs, such as, for example, by elimination of an alcohol derivative such as a xanthate ester, or a halide. Alternatively the unsaturation can be present in one of the precursors for the chain. For example an unsaturated fatty acid could be converted to an ω -amino unsaturated acid, for example, via the ω -bromo derivative. This material could then be used to form the ring as described for the saturated analogs. It should be noted that compounds containing unsaturation in the ring are amenable to further transformation. Cycloadditions such as Diels-Alder reactions, 1,3-dipolar additions, and 2+2 cycloadditions lead to bridged 6-, 5-, and 4-membered ring systems. Three membered rings can be generated by carbene reactions, epoxidations with, for example, peracids, and aziridinations. Alternatively the ring can be part of the ω -amino acid before attachment to the core. Thus, for example, ω -aminoalkyl benzoic acids could be utilized to form the chain.

Carbamates, thiocarbamates and ureas at Y may be generated by reacting the C-protected amino acid core with a phosgene equivalent such as trichloroacetyl chloride followed by a mono N-protected ω -diamine, ω -amino alcohol,

or ω -amino thiol. Similarly sulfamides can be generated by replacing the phosgene equivalent by sulfonyl chloride, or an equivalent.

Compounds of formula IVa may be generated by replacing the amino acid core in IV by a diamine or equivalent. One way this can be accomplished is by taking a suitably N-protected amino acid, forming an active ester, displacing by NaN_3 to form the acyl azide, and then heating to form the isocyanate. Reaction with mono N-protected ω -diamines, ω -amino alcohols, or ω -amino thiols as above provides carbamates, thiocarbamates and ureas at X. Alternatively, the isocyanate can be hydrolyzed to give the amine which can be acylated to form amide derivatives. Similarly, sulfonylation can provide sulfonyl analogs.

Compounds of formula IVb replace the amine at X in IVa by oxygen or sulfur. These compounds can be synthesized, for example, by using phenylalaninal as the starting material. Conversion of the aldehyde to the epoxide may be accomplished either directly with an appropriate ylide, or indirectly via Wittig olefination followed by epoxidation, for example, with a peracid. The epoxide can then be reacted with a suitable alcohol or thiol to provide a diol or mercapto alcohol respectively. Acylation of these moieties, with for example a suitable isocyanate provides the X functionality described in IVb.

Compounds of formula V may be prepared in a similar fashion as those for formula IV but using an optionally α -alkylated homostatine core rather than a statine core. Homostatine analogs can be synthesized, for example, by activating the carboxylic acid of a protected statine analog, reacting with diazomethane to generate the diazoketone, and then subjecting it to Wolff rearrangement using thermal, photochemical, or metal-catalyzed conditions. The resulting carboxylic acid can then be elaborated as in formula IV.

Compounds of formula Va have the amide of X in formula V reversed. Such compounds are available from derivatives of statine. Thus a statinal derivative can be reductively aminated to give the required precursor amine which can then be acylated, sulfonylated, or phosphorylated as for previously described examples.

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Alternatively the amine could be synthesized by reduction of a statine amide or reaction of an activated statinol derivative with an amine equivalent.

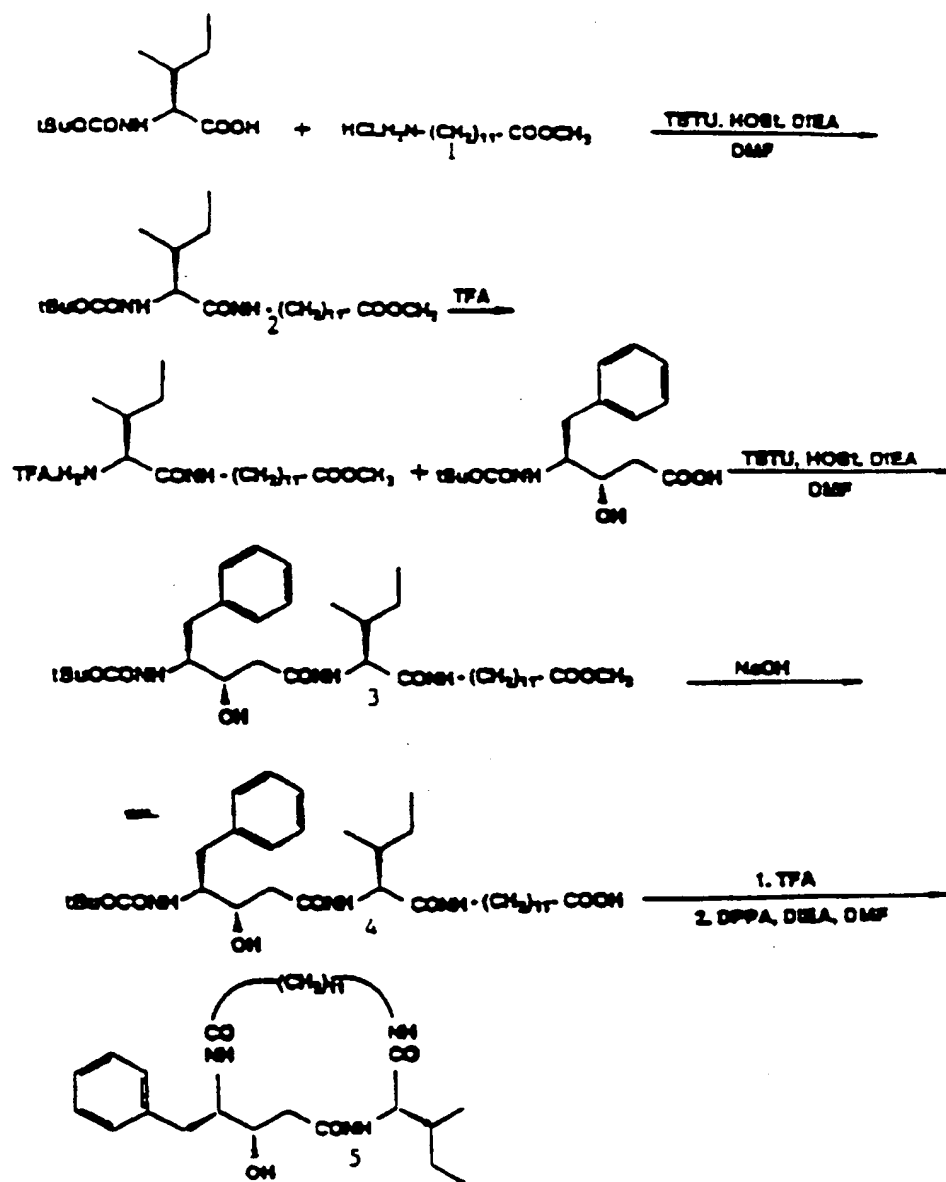
Similarly, compounds of formula Vb can be derived from statinol analogs by acylation on oxygen with, for example, an isocyanate derivative. Alternatively, the alcohol can be activated as, for example, a sulfonate derivative and then reacted with a thiol to give the sulfur analogs.

The Examples herein are meant to exemplify the various aspects of carrying out the invention and are not intended to limit the scope of the invention in any way.

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EXAMPLE 1

The synthesis of this inhibitor is represented in Scheme A below:



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The various compounds shown in scheme A can be prepared as follows:

12-aminododecanoic acid methylester hydrochloride (1)

To the suspension of 12-aminododecanoic acid (2.15 g; 10 mmol) in 2,2-dimethoxypropane (100 ml) was added 36% aqueous HCl (10 ml) and the mixture was treated in an ultrasonic bath until all solid material was dissolved. The mixture was then left to stand overnight, concentrated to dryness, dissolved in 40 ml of methanol and precipitated with approximately 500 ml of dry ether to give 2.4 g of 1 (90.5% yield).

12-(t-Butyloxycarbonyl-isoleucinyl)-aminododecanoic acid methylester (2)

Boc-Ile-NH-(CH₂)₁₁-COOCH₃

Boc (t-butyloxycarbonyl)-Ile (isoleucine)-OH. 1/2H₂O (384 mg; 1.6 mmol), HCl.H₂N-(CH₂)₁₁-COOCH₃ (1) (398 mg; 1.5 mmol), HOBT (N-hydroxybenzotriazole) (260 mg; 1.7 mmol) and TBTU (2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate) (546 mg; 1.7 mmol) were dissolved in DMF (5 ml). DIEA (diisopropylethylamine) (860 µl; 5 mmol) was added to the above solution and the mixture was left to stand overnight. Five drops of N-aminoethylmorpholine were then added to the solution. After 30 minutes DMF was removed in vacuo and the solid residue suspended in 10% aqueous KHSO₄ (20 ml). The precipitate was filtered off, washed with water, 5% aqueous NaHCO₃, water again, and then dried in a vacuum desiccator to give 630 mg of 2 (95% yield).

12-(t-Butyloxycarbonyl-phenylstatinyl-isoleucinyl)-aminododecanoic acid methylester (3)

Boc-Pst-Ile-NH-(CH₂)₁₁-COOCH₃

Boc-Ile-(CH₂)₁₁-COOCH₃ (2) (384 mg; 0.8 mmol) was dissolved in TFA (trifluoroacetic acid) (3 ml). After 30 minutes of standing at room temperature, the TFA was removed in vacuo, the oily residue was concentrated three more times with ether to remove the remaining TFA, and then dissolved in 5 ml of DMF. Boc-Pst-OH

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(247 mg; 0.8 mmol), TBTU (289 mg; 0.9 mmol) and DIEA (1030 μ l; 6 mmol) were added to the above solution and the mixture was left to stand overnight. Workup as for 2 yielded 492 mg of 3 (97%).

12-(t-Butyloxycarbonyl-phenylstatinyl-isoleucinyl) aminododecanoic acid (4)

Boc-Pst-Ile-NH-(CH₂)₁₁-COOH

Boc-Pst-Ile-NH-(CH₂)₁₁-COOCH₃ (3) (380 mg; 0.6 mmol) was dissolved in a mixture of 1,4 dioxane (5 ml) and methanol (1 ml) and combined with 1N aqueous solution of NaOH (1.5 ml). The reaction was complete in approximately two hours. The organic solvents were evaporated and the remaining aqueous slurry was acidified with 10% aqueous KHSO₄ (10 ml). The precipitate was filtered off, washed with water, and dried in a vacuum desiccator to yield 353 mg of 4 (95% yield).

Cyclo-12-aminododecanoyl-phenylstatinyl-isoleucinyl (5)

-CO-Pst-Ile-NH-(CH₂)₁₁-

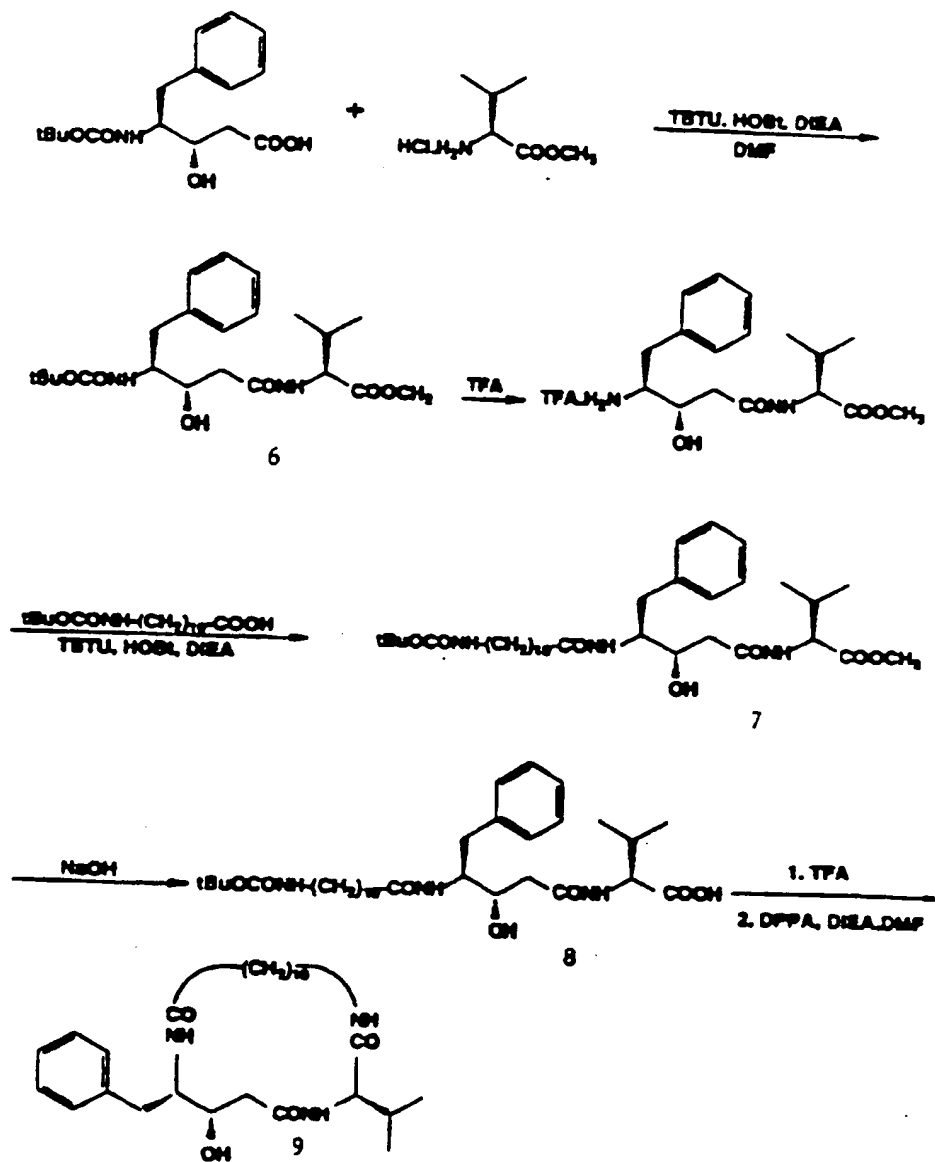
Boc-Pst-Ile-NH-(CH₂)₁₁-COOH (4) (248 mg; 0.4 mmol) was deprotected with TFA as described for 3 and dissolved in DMF (20 ml). This solution was added dropwise to a stirred solution of DPPA (diphenyl phosphorylazide) (1.1 g; 4 mmol) and DIEA (1.38 ml; 8 mmol) in DMF (100 ml) over the course of 17 hours via syringe pump. The mixture was then stirred for an additional 3 hours, evaporated to dryness and triturated with ether. The precipitated product was filtered off, washed with ether, 5% aqueous NaHCO₃, and water and then dried in a vacuum desiccator to provide 160 mg of crude 5 (80% yield). For the purpose of biological testing, the product was purified by RP-HPLC.

SIMS MS M⁺Na⁺524

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EXAMPLE 2

An alternate sequence to a cyclic inhibitor is represented in Scheme B below:

SCHEME B:

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t-Butyloxycarbonyl-phenylstatinyl valine methylester (6)Boc-Pst-Val-OCH₃

Boc-Pst-OH (309 mg; 1 mmol), HCl.H-ValOCH₃ (184 mg; 1.1 mmol) and TBTU (385 mg; 1.2 mmol) were dissolved in 5 ml of DMF and DIEA (688 μ l; 4 mmol) was added to the solution. The mixture was left to stand overnight and then evaporated to dryness. The oily residue was dissolved in ethylacetate and the ethylacetate solution was washed with 10% aqueous KHSO₄, water, 5% aqueous NaHCO₃, and water again, dried over MgSO₄, and concentrated to give 420 mg of 6.

11-t-Butyloxycarbonyl-aminoundecanoyl-phenylstatinyl valine methyl ester (7)Boc-NH-(CH₂)₁₀-CO-Pst-Val-OCH₃

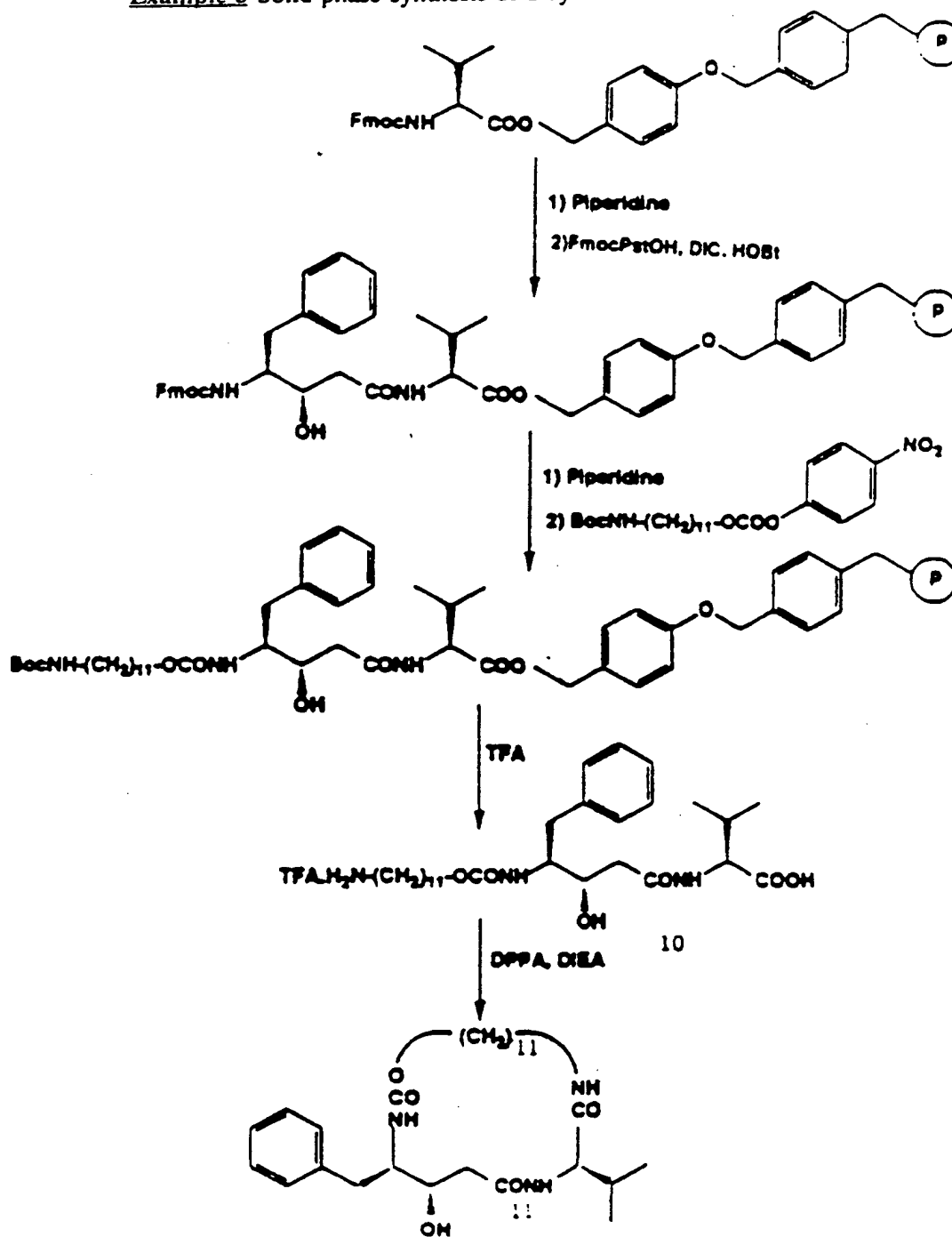
Boc-Pst-Val-OCH₃ (420 mg; 1 mmol) was deprotected with TFA as described for 3. After addition of ether, a precipitate of TFA.H-Pst-Val-OCH₃ (349 mg; 0.8 mmol) was obtained. This was dissolved in 5 ml of DMF together with Boc-NH-(CH₂)₁₀-COOH (241 mg; 0.8 mmol), TBTU (289 mg; 0.9 mmol) and DIEA (516 μ l; 3 mmol). After 10 hours of standing at room temperature, DMF was removed in vacuo, the residue dissolved in ethylacetate and worked up as described above for 6 to yield 480 mg of oily 7.

11-t-Butyloxycarbonyl-aminoundecanoyl-phenylstatinyl valine (8)Boc-NH-(CH₂)₁₀-CO-Pst-ValOH

Hydrolysis of crude Boc-NH(CH₂)₁₀-CO-Pst-ValOCH₃ (7) (480 mg; 0.8 mmol) as described above for 4 provided 435 mg of oily 8.

Cyclo-11-aminoundecanoyl-phenylstatinyl-valine (9)-CO-Pst-Val-NH-(CH₂)₁₀-

Boc-NH-(CH₂)₁₀-CO-Pst-ValOH (8) was deprotected and cyclized using the procedure described for 5. Yield 260 mg (55% from Boc-Pst-OH used in 6) of crude solid 9. For the purpose of biological testing the product 9 was purified by RP-HPLC. SIMS MS MH⁺474

Example 3 Solid phase synthesis of a cyclic inhibitor.

(SEQ.ID.NO.48)

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Solid Phase Synthesis of TFA.H₂N-(CH₂)₁₁-OCOPstVal-OH (10)

FmocVal-WANG-resin (1g, 0.37 mmol) was swollen in DMF, washed with DMF (3x2 min), deprotected with 20% piperidine in DMF (1x2 min; 1x20 min) and washed again with DMF (5x2 min). FmocPstOH (319 mg, 0.74 mmol) was then coupled using DIC (diisopropylcarbodiimide) (101 mg, 0.8 mmol) and HOBt (122 mg, 0.8 mmol) for activation. The resulting resin was then deprotected and washed as described above. The solution of BocNH-(CH₂)₁₁-COO-pC₆H₄-NO₂ (452 mg, 1 mmol) and DIEA (190 μ l, 1.1 mmol) in 5 ml of DMF was added and the mixture was shaken overnight. The resin was washed with DMF (5x2 min), 20% piperidine in DMF (2x2 min), DMF (3x2 min), pyridine-DCM (1:1) (3x2 min) and DCM (dichloromethane) (5x2 min). The product was then cleaved off the resin with 95% TFA (2x30 min) and the resin was washed with DCM (5x2 min). The combined solutions were evaporated to dryness. The oily residue (200 mg) was used directly for the next reaction.

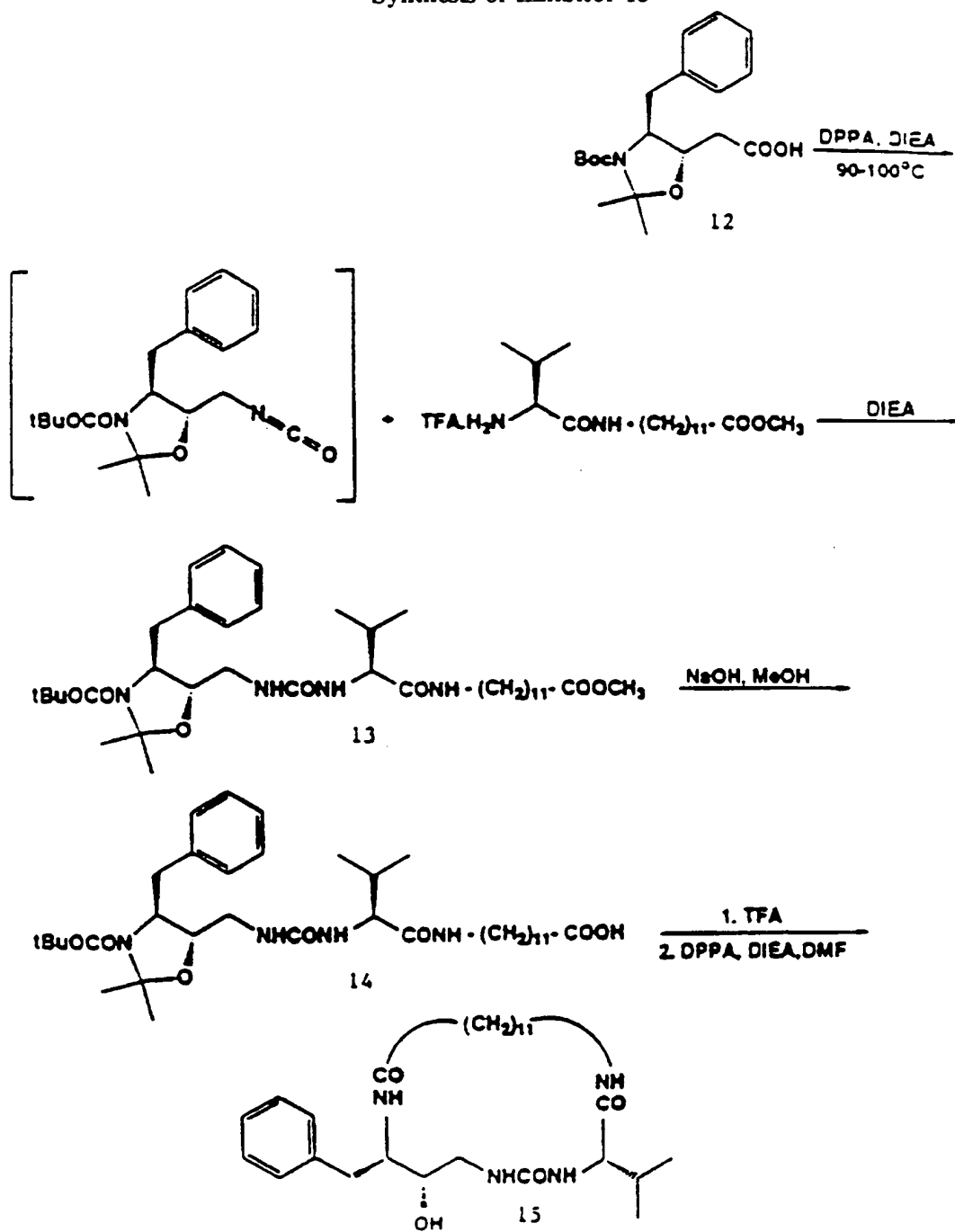
Synthesis of (11) (cyclization)

TFA.H₂N-(CH₂)₁₁-OCOPstVal-OH(10) was cyclized using the procedure described for 5. Yield 40 mg (21.5% based on the resin substitution) of crude 11. For the purpose of biological testing the product was purified by RP-HPLC. Product was characterized by SIMS-MS, M+H⁺ = 504, M+Na⁺ = 526. Compound 11 has a K_i against cathepsin D of 3.6 nM.

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Example 4 Synthesis of a cyclic urea.

Synthesis of Inhibitor 15



(SEQ.ID.NO.49)

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Synthesis of 13

Boc-Pst(acetonide)OH (850 mg, 2.44 mmol) was dissolved in 13 ml of toluene and approx. 2-3 ml of toluene was distilled off to dry the solution. DPPA (1 g, 3.66 mmol) and DIEA (0.84 ml, 4.88 mmol) were added and the mixture was heated to 90-100°C for 2 hours. A solution of TFA.H-ValNH-(CH₂)₁₁-COOMe prepared from 1.2 g (2.8 mmol) Boc-ValNH(CH₂)₁₁CO₂Me and DIEA (1.46 ml, 8.5 mmol) in 3 ml of dry toluene was added to the reaction mixture, the mixture was heated for an additional 30 min and then left to stand overnight at room temperature. The toluene was removed in vacuo, the residue dissolved in ethyl acetate and washed subsequently with 10% aq. KHSO₄, brine, 5% NaHCO₃, brine, and finally dried with MgSO₄. The ethyl acetate was then removed in vacuo and the oily residue purified by flash chromatography in hexane-ethyl acetate (3:2). Yield 873 mg, 53% of 13. Product was characterized by SIMS-MS, M+H⁺ = 675.

Synthesis of 14

13 (850 mg, 1.26 mmol) was hydrolyzed with NaOH according to the procedure described above for 4. Yield 850 mg (~ 100%) of oily 14, that was used directly for the next reaction.

Synthesis of 15 (Cyclization)

This compound was prepared using the procedure described above for 5. Yield 438 mg (69% calculated from 13). Product was characterized by SIMS-MS, M+H⁺ = 503. Compound 15 has a K_i against cathepsin D of 45 nM.

Inhibition Assays

Kinetic measurements. Fluorogenic substrates Ac-Glu-Glu(EDANS)-Lys-Pro-Ile-Cys-Phe-Phe-Arg-Leu-Gly-Lys(DABCYL)-Glu-NH₂ and Ac-Glu-Glu(EDANS)-Lys-Pro-Ile-Cys-Phe-Leu-Arg-Leu-Gly-Lys(DABCYL)-Glu-NH₂ were used for measuring the activity of cathepsin D and plasmepsin 2 correspondingly. Typically, 485 µl of

50 mM Gly-HCl buffer, pH 3.5 (in the case of cathepsin D), or 100 mM sodium acetate buffer, pH 5.0 (in the case of plasmepsin 2), was mixed with 5 μ l of DMSO and 5 μ l of titrated protease (final concentration 0.2-10 nM) and incubated 3 min at 37°C. The reaction was initiated by the addition of 5 μ l of substrate stock solution in DMSO. Increase in fluorescence intensity at the emission maximum of 487 nm (excitation wavelength was 349 nm) was monitored as a function of time using an Aminco Bowman-2 luminescence spectrometer (SLM Instruments, Inc.).

Plasmepsin 1 assays were run similarly to plasmepsin 2, using the fluorogenic substrate DABCYL-Gaba-Glu-Arg-Met-Phe-Leu-Ser-Pro-Gaba-Glu(EDANS)-NH₂.

The initial rate of hydrolysis was calculated by a second degree polynomial fit using SLM AB2 2.0 operating software. Kinetic parameters were determined by nonlinear regression fitting of initial rate versus substrate concentration data to the Michaelis-Menten equation using the program Enzfitter version 1.05 (Leatherbarrow, R.J. 1987. Enzfitter, a program for non-linear regression analysis. Elsevier Scientific, New York).

For inhibition studies inhibitors were prepared as stock solutions at different concentrations in DMSO. In a typical experiment 485 μ l of the appropriate buffer was mixed with 5 μ l of inhibitor stock solution and 5 μ l of titrated protease (final concentration 0.2-10 nM) and preincubated 3 min at 37°C. The reaction was initiated by the addition of 5 μ l of substrate stock solution in DMSO. For data analysis the mathematical model for tight-binding inhibitors (Williams, J.W., and Morrison, J.F. Methods Enzymol. 63: 437 (1979)) was used. The data were fitted by nonlinear regression analysis to the equation

$$V = V_0 / 2E_t \left(\{ [K_i(1 + S/K_m) + I_t - E_t]^2 + 4K_i(1 + S/K_m)E_t \}^{1/2} - [K_i(1 + S/K_m) + I_t - E_t] \right)$$

with the program Enzfitter (version 1.05), where V and V₀ are initial velocities with and without inhibitor, K_m is a Michaelis-Menten constant and S, E_t and I_t are the concentrations of substrate, active enzyme and inhibitor respectively.

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Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it will be obvious to those skilled in the art that certain changes and modifications may be practiced without departing from the spirit and scope thereof as described in the specification and as defined in the appended claims.

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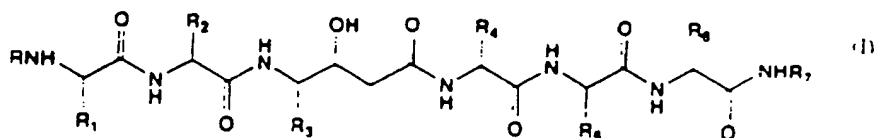
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WHAT IS CLAIMED IS:

1. A compound of formula (I):



in which R, R₇ = represent hydrogen, alkoxycarbonyl, aralkoxy-carbonyl, alkylcarbonyl, cycloalkylcarbonyl, cycloalkylalkoxy-carbonyl, cycloalkylalkanoyl, aralkanoyl, aroyl, aryloxycarbonyl, aryloxycarbonylalkyl, aryloxyalkanoyl, heterocyclylcarbonyl, hetero-cyclyloxycarbonyl, heterocyclylalkanoyl, heterocyclylalkoxycarbonyl, heteroaralkanoyl, heteroaralkoxycarbonyl, heteroaryloxy-carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxyalkyl, heteroaryloxyalkyl, hydroxyalkyl, aminocarbonyl, aminoalkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted aminoalkyl, alkoxy-alkyl, alkylthioalkyl, mono- and disubstituted aminoalkanoyl radicals wherein the substituents are selected from alkyl, aryl, aralkyl, cycloalkyl, cyclo-alkylalkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkylalkyl radicals, or where said aminoalkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

R₁, R₂, R₄, R₅, R₆ = optionally substituted lower alkyl, lower cycloalkyl, aryl, aralkyl and heteroaryl;

R₃ = optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, aralkyl, aralkoxy, aralkylthio, aralkylamino, cycloalkyl-alkyl, cycloalkylalkoxy, cycloalkylalkylthio, cycloalkylalkylamino.

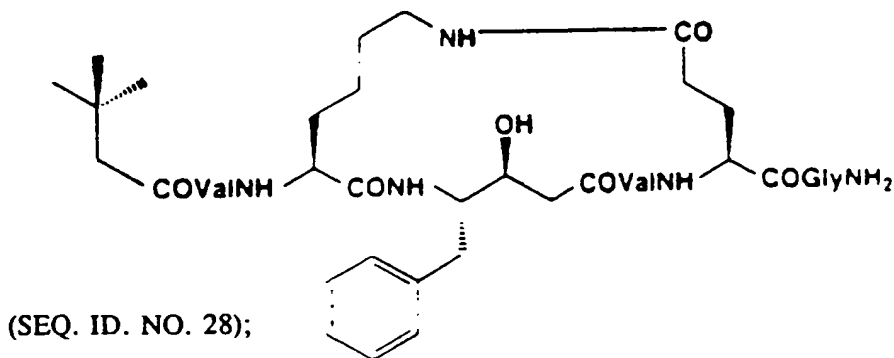
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2. The compound of claim 1, wherein R_2 and R_3 or R_1 and R_3 are connected by an optionally substituted bridging moiety comprised of a stable combination of C, N, O or S atoms.

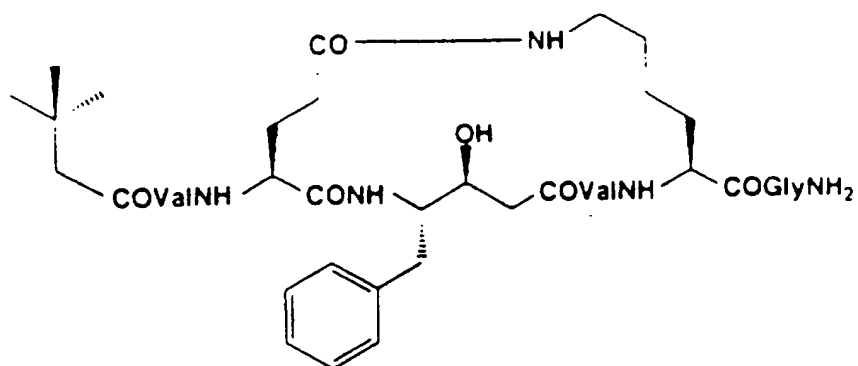
3. The compound of claim 1, wherein R_1 , R_2 , R_4 , R_5 and R_6 represent the side chain or a residue of an amino acid after formation of a peptide linkage.

4. The compound of claim 1, which is selected from the group consisting of

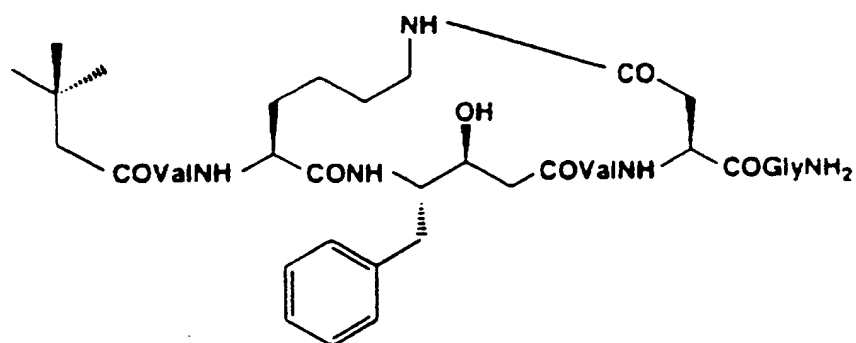
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- (2) Iva-Gln-Val-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 2);
- (3) Iva-Lys-Val-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 3);
- (4) Tba-Val-Val-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 4);
- (5) Iva-Val-Ile-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 5);
- (6) Iva-Val-Leu-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 6);
- (7) Tba-Val-Val-Pst-Val-Leu-Gly-NH₂ (SEQ. ID. NO. 7);
- (8) Tba-Val-Cys-Pst-Val-Leu-Gly-NH₂ (SEQ. ID. NO. 8);
- (9) Tba-Val-Glu-Pst-Val-Leu-Gly-NH₂ (SEQ. ID. NO. 9);
- (10) Tba-Val-Asp-Pst-Val-Leu-Gly-NH₂ (SEQ. ID. NO. 10);
- (11) Iva-Val-Val-Sta-Ala-Leu-Gly-NH₂ (SEQ. ID. NO. 11);
- (12) Tba-Val-Cys-Pst-Val-Cys-Gly-NH₂ (SEQ. ID. NO. 12).



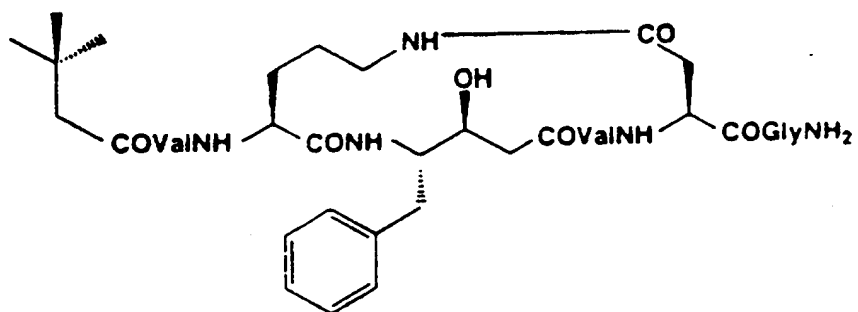
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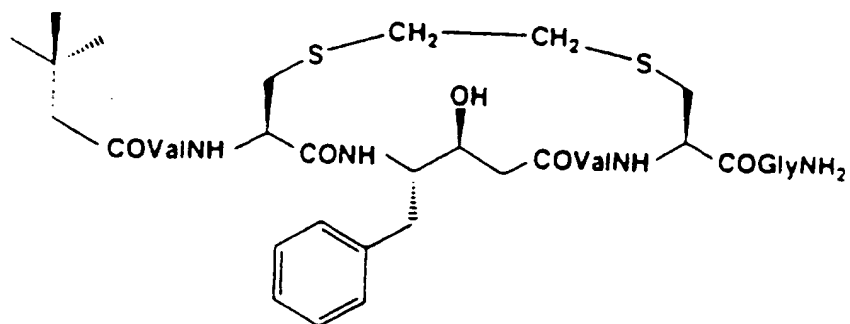


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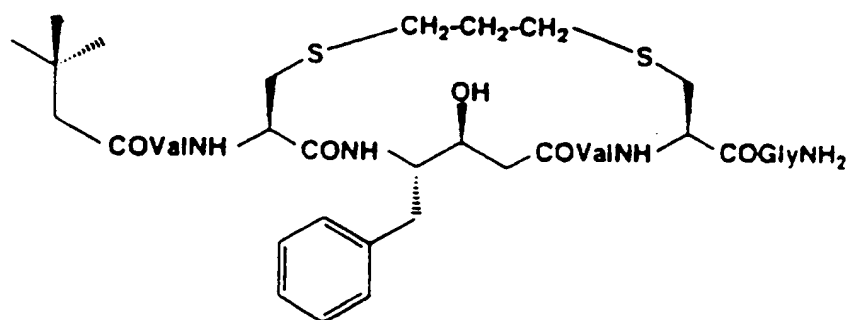


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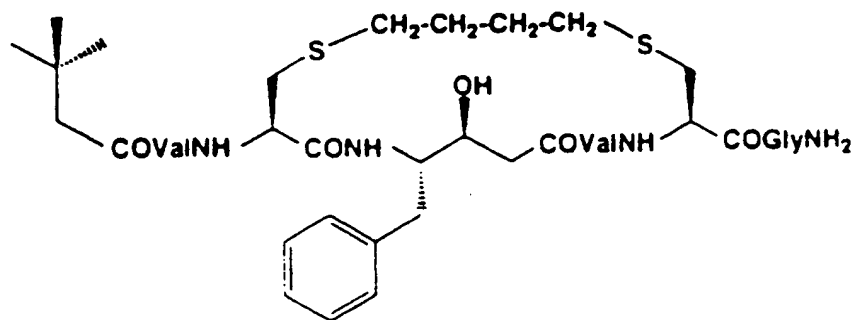
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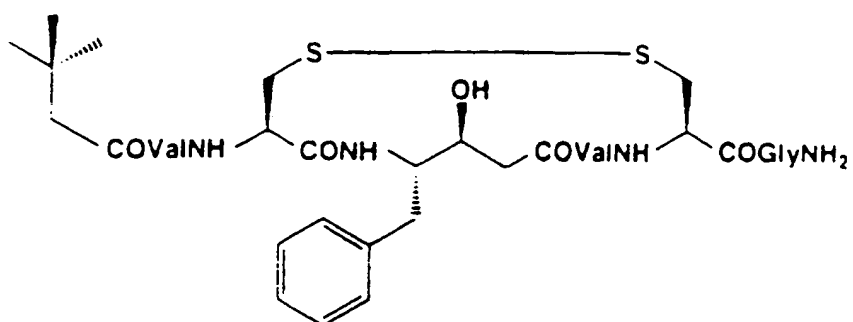


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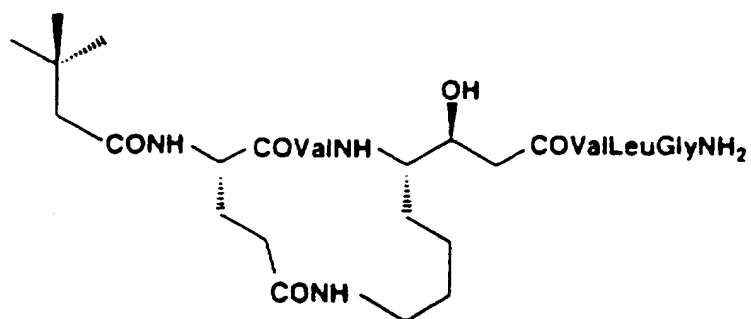


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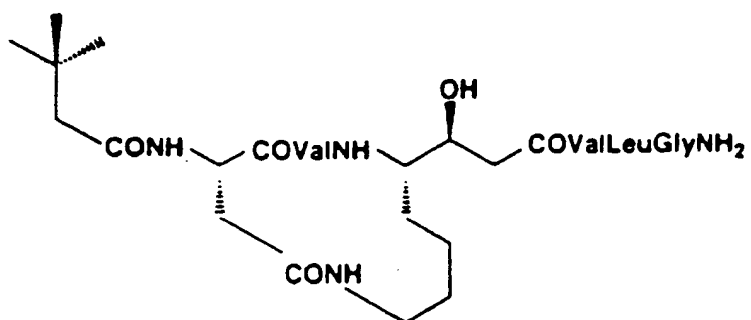
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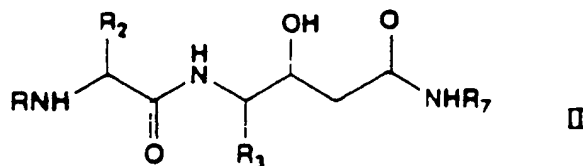
(SEQ. ID. NO. 36);



(SEQ. ID. NO. 37).

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5. A compound of formula II



in which R, R₇ = represent hydrogen, alkoxycarbonyl, aralkoxycarbonyl, alkylcarbonyl, cycloalkylcarbonyl, cycloalkylalkoxycarbonyl, cycloalkylalkanoyl, aralkanoyl, aroyl, aryloxy-carbonyl, aryloxycarbonylalkyl, aryloxyalkanoyl, heterocyclylcarbonyl, heterocyclyloxycarbonyl, heterocyclylalkanoyl, heterocyclylalkoxycarbonyl, heteroaralkanoyl, heteroaralkoxycarbonyl, heteroaryloxy-carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxyalkyl, heteroaryloxyalkyl, hydroxyalkyl, aminocarbonyl, aminoalkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted aminoalkyl, alkoxyalkyl, alkylthioalkyl, mono- and disubstituted aminoalkanoyl radicals wherein the substituents are selected from alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkylalkyl radicals, or where said aminoalkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

R₂ = optionally substituted lower alkyl, lower cycloalkyl, aryl, aralkyl and heteroaryl;

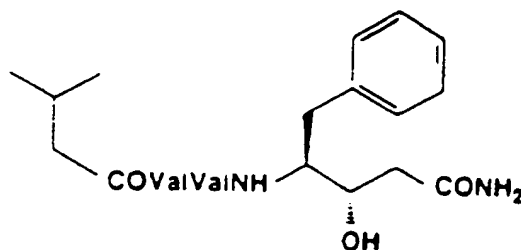
R₃ = optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, aralkyl, aralkoxy, aralkylthio, aralkylamino, cycloalkylalkyl, cycloalkylalkoxy, cycloalkylalkylthio, cycloalkylalkylamino.

6. The compound of claim 5, wherein R and R₃ are connected by an optionally substituted bridging moiety comprised of a stable combination of C, N, O or S atoms.

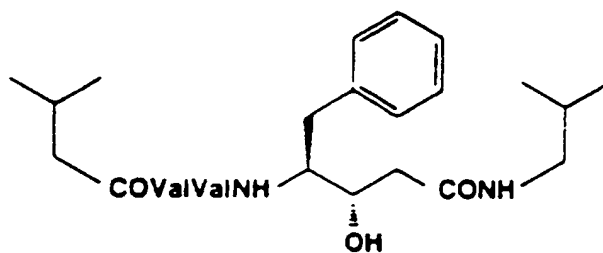
- 53 -

7. The compound of claim 5, wherein R_2 represents the side chain of a residue of an amino acid after formation of a peptide linkage.

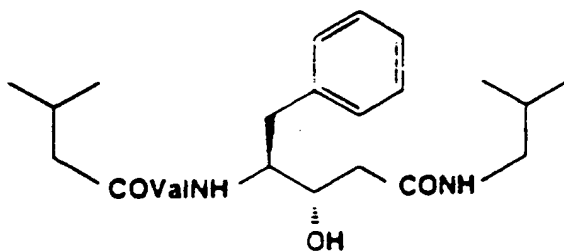
8. The compound of claim 5, which is selected from the group consisting of



(SEQ. ID. NO. 13);

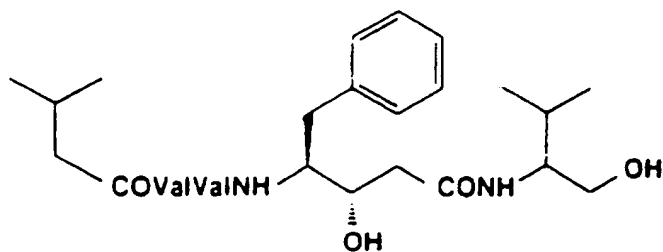


(SEQ. ID. NO. 14);

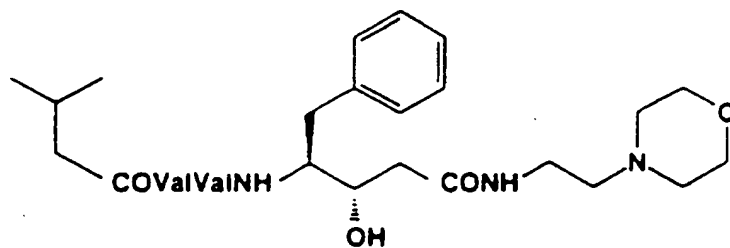


(SEQ. ID. NO. 15);

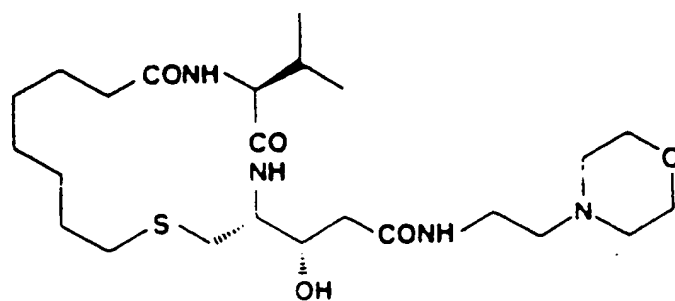
- 54 -



(SEQ. ID. NO. 16);



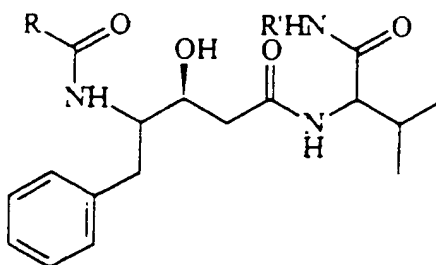
(SEQ.ID.NO. 17)



(SEQ.ID.NO. 38)

- 55 -

9. The compound of claim 5, having the formula:



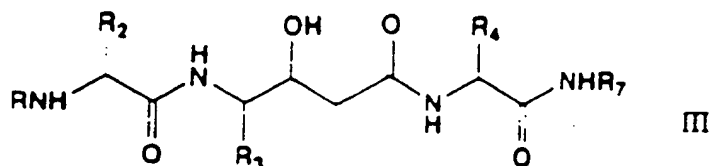
in which R and R¹ are given in the following table:

R	R ¹
(CH ₂) ₄ CH ₃	(CH ₂) ₆ CH ₃
(CH ₂) ₂ SPr	(CH ₂) ₂ SPr
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₂ SCH ₂ Ph
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₃ SCH ₂ Ph
(CH ₂) ₃ SCH ₂ Ph	(CH ₂) ₃ SCH ₂ Ph
O(CH ₂) ₃ -4-Pyr	(CH ₂) ₃ SCH ₂ Ph
(CH ₂) ₃ SCH ₂ Ph	(CH ₂) ₆ CH ₃
(CH ₂) ₂ SCH ₂ Ph	(CH ₂) ₆ CH ₃
(CH ₂) ₄ CH ₃	(CH ₂) ₃ SBn
(CH ₂) ₂ CO ₂ H	n-C ₇ H ₁₅
(CH ₂) ₂ CO ₂ Me	n-C ₇ H ₁₅
(CH ₂) ₂ COHN ₂	n-C ₇ H ₁₅
(CH ₂) ₃ CO ₂ H	n-C ₇ H ₁₅
(CH ₂) ₃ CO ₂ Me	n-C ₇ H ₁₅
CH ₂ -p-Ph-Ph	n-C ₇ H ₁₅
(E)-3-	n-C ₇ H ₁₅

CH=CHbenzodioxolane

Ph = phenyl, pyr = pyridyl and Bn = benzyl.

10. A compound of formula III:



in which R, R₇ = represent hydrogen, alkoxycarbonyl, aralkoxycarbonyl, alkylcarbonyl, cycloalkylcarbonyl, cycloalkylalkoxycarbonyl, cycloalkylalkanoyl, aralkanoyl, aroyl, aryloxycarbonyl, aryloxycarbonylalkyl, aryloxyalkanoyl, heterocyclylcarbonyl, heterocyclyloxycarbonyl, heterocyclylalkanoyl, heterocyclylalkoxycarbonyl, heteroaralkanoyl, heteroaralkoxycarbonyl, heteroaryloxy-carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxyalkyl, heteroaryloxyalkyl, hydroxyalkyl, aminocarbonyl, aminoalkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted aminoalkyl, alkoxyalkyl, alkylthioalkyl, mono- and disubstituted aminoalkanoyl radicals wherein the substituents are selected from alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkylalkyl radicals, or where said aminoalkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

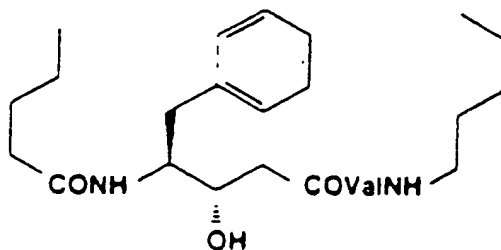
R₂, R₄ = optionally substituted lower alkyl, lower cycloalkyl, aryl, aralkyl and heteroaryl;

R₃ = optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, aralkyl, aralkoxy, aralkylthio, aralkylamino, cycloalkylalkyl, cycloalkylalkoxy, cycloalkylalkylthio, cycloalkylalkylamino.

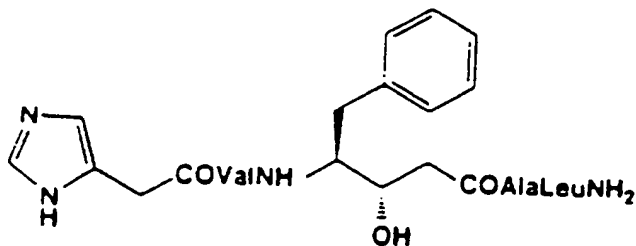
11. The compound of claim 10, wherein R₂ and R₄ represent side chains of a residue of an amino acid after formation of a peptide linkage.

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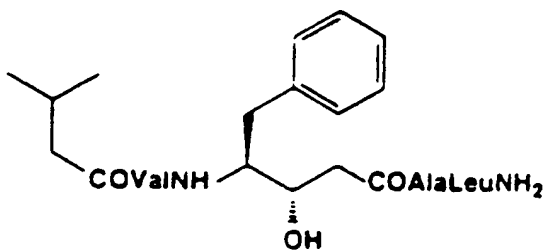
12. The compound of claim 10, which is selected from the group consisting of



(SEQ. ID. NO. 18);

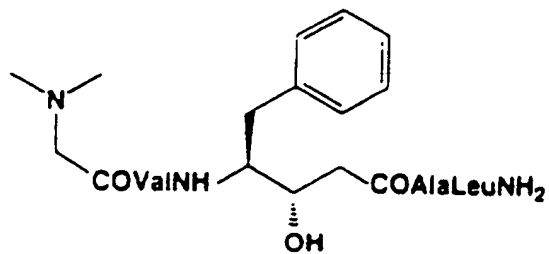


(SEQ. ID. NO. 19);

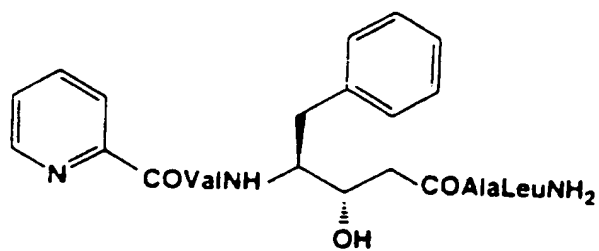


(SEQ. ID. NO. 20);

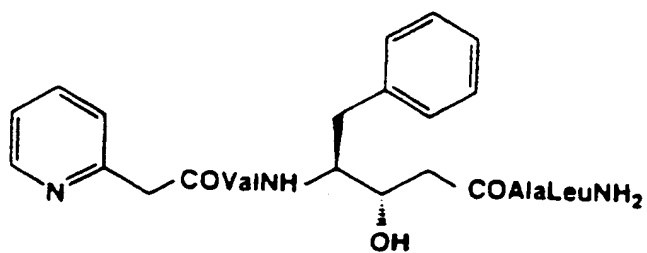
- 58 -



(SEQ. ID. NO. 21);

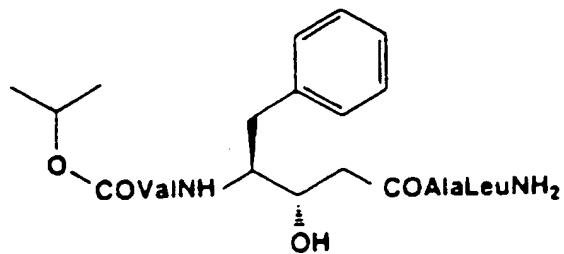


(SEQ. ID. NO. 22);

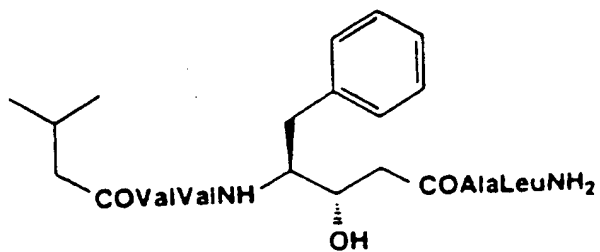


(SEQ. ID. NO. 23);

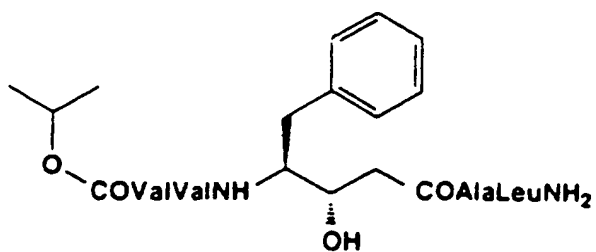
- 59 -



(SEQ. ID. NO. 24);

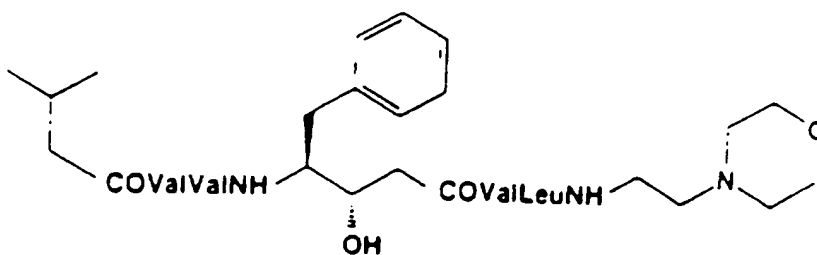


(SEQ. ID. NO. 25);



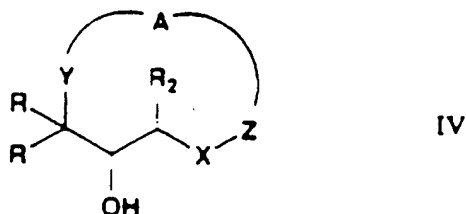
(SEQ. ID. NO. 26);

- 60 -



(SEQ. ID. NO. 27).

13. A compound of the formula IV:



where $X = \text{CWNR}, \text{CWOR}, \text{S(O)}_n\text{NR}, \text{P(O)(Q)NR}$; $W = \text{O}, \text{S}, \text{NR}$; $n = 0, 1, 2$;

$Q = \text{R}, \text{OR}, \text{N(R)}_2$

$Y = \text{NRCW}, \text{NRS(O)}_n, \text{NRP(O)(Q)}$

$Z = \text{CRR}', \text{NR}$

A = an optionally substituted bridging group which is saturated or unsaturated and having 2-15 atoms comprised of any stable combination of C, N, O, or S.

R = represents hydrogen, alkoxycarbonyl, aralkoxycarbonyl, alkylcarbonyl, cycloalkylcarbonyl, cycloalkylalkoxycarbonyl, cycloalkylalkanoyl, aralkanoyl, aroyl, aryloxycarbonyl, aryloxycarbonylalkyl, aryloxyalkanoyl, heterocyclcarbonyl, heterocyclloxycarbonyl, heterocycllalkanoyl, heterocycllalkoxy-carbonyl, heteroaralkanoyl, heteroaralkoxycarbonyl, heteroaryloxy-carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxyalkyl, hetero-aryloxyalkyl, hydroxyalkyl, aminocarbonyl, aminoalkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted aminoalkyl, alkoxyalkyl, alkylthioalkyl, mono- and disubstituted aminoalkanoyl radicals wherein the substituents are

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selected from alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkylalkyl radicals, or where said aminoalkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

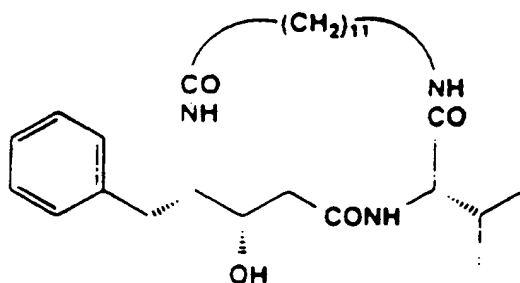
$R_1, R_2 = H$, halo, optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino;

$R_3 =$ optionally substituted lower alkyl, lower alkoxy, lower alkylthio, mono or di-lower alkyl amino, aralkyl, aralkoxy, aralkylthio, aralkylamino, cycloalkylalkyl, cycloalkylalkoxy, cycloalkylalkylthio, cycloalkylalkylamino.

14. The compound of claim 13, wherein $X, Y = NRCW, NRS(O)_n, NRP(O)(Q)$ and; $Z = CRR', NR, O, S$.

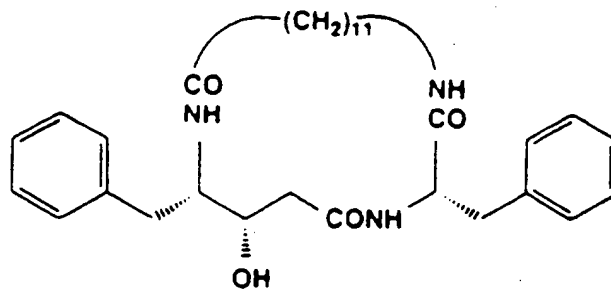
15. The compound of claim 13, wherein $X = OCWNR, SCWNR$; $Y = NRCW, NRS(O)_n, NRP(O)(Q)$ and; $Z = CRR', NR$.

16. The compounds of claim 13, which are selected from the group consisting of

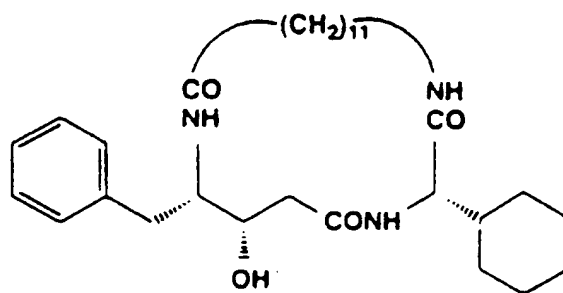


(SEQ.ID.NO. 39);

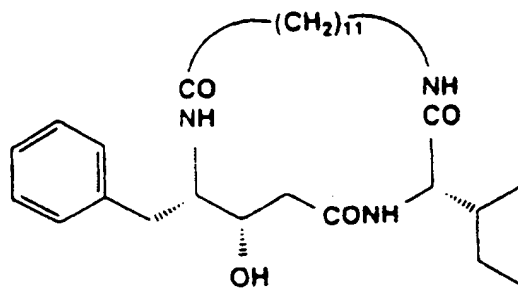
- 62 -



(SEQ.ID.NO. 40);

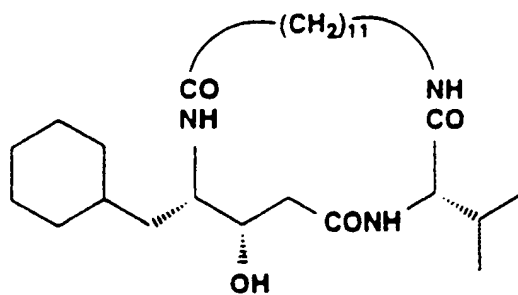


(SEQ.ID.NO. 41);

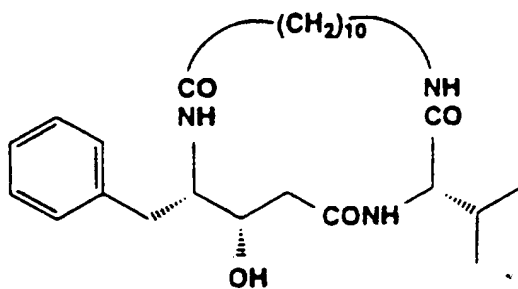


(SEQ.ID.NO. 42);

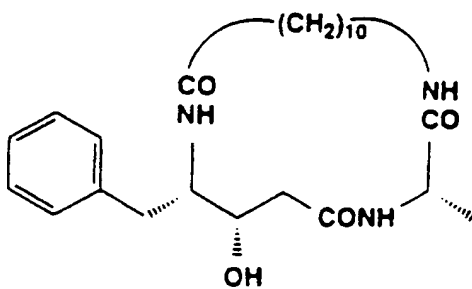
- 63 -



(SEQ.ID.NO. 43);

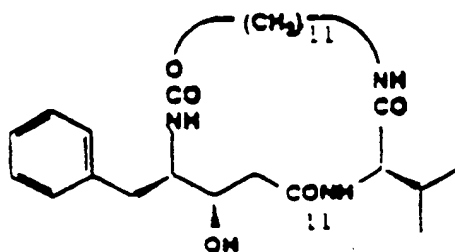


(SEQ.ID.NO. 44);

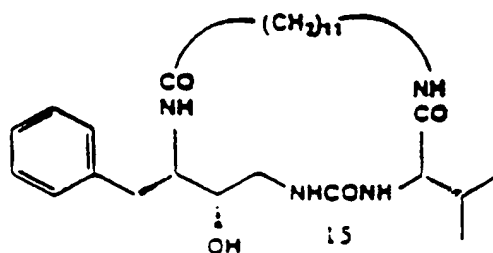


(SEQ.ID.NO. 45)

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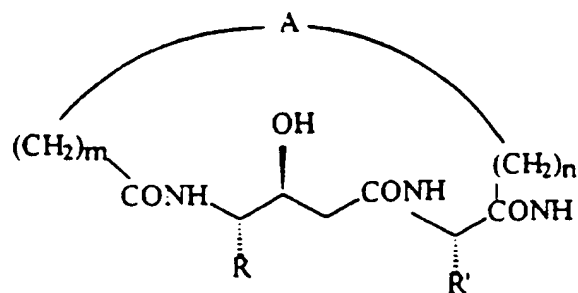


(SEQ.ID.NO. 48)



(SEQ.ID.NO. 49)

17. The compound of claim 13, having the formula:



in which R and R' are given in the following table:

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R	R ⁱ	m	n	A
PhCH ₂	D-iPr	11	-	-
PhCH ₂	iBu	11	-	-
PhCH ₂	Me	11	-	-
PhCH ₂	Et	11	-	-
PhCH ₂	3-PyrCH ₂	11	-	-
PhCH ₂	CH ₃ CH(OCH ₂ Ph) CH ₂	11	-	-
PhCH ₂	CH ₃ CH(OH)CH ₂	11	-	-
PhCH ₂	tBu	11	-	-
PhCH ₂	Ph	11	-	-
p-(PhCH ₂ O)- PhCH ₂	iPr	11	-	-
p-(OH)PhCH ₂	iPr	11	-	-
p- (OMe)PhCH ₂	iPr	11	-	-
p-(OEt)- PhCH ₂	iPr	11	-	-
p-(OiPr)- PhCH ₂	iPr	11	-	-
p-(3PyrCH ₂ O)- PhCH ₂	iPr	11	-	-

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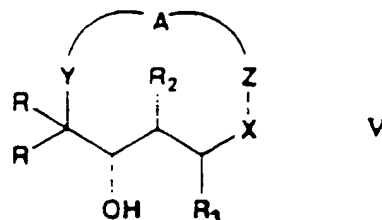
R	R ¹	m	n	A
p- MeO(CH ₂) ₂ O- PhCH ₂	iPr	11	-	-
(F ₃ -Ph)CH ₂	iPr	11	-	-
(m,p- Cl ₂ Ph)CH ₂	iPr	11	-	-
(mp-F ₂ Ph)CH ₂	iPr	11	-	-
p-F-PhCH ₂	iPr	11	-	-
H SCH ₂	iPr	11	-	-
MeSCH ₂	iPr	11	-	-
EtSCH ₂	iPr	11	-	-
PrSCH ₂	iPr	11	-	-
CH ₂ =CHCH ₂ SCH ₂	iPr	11	-	-
BuSCH ₂	iPr	11	-	-
iBuSCH ₂	iPr	11	-	-
PhCH ₂ SCH ₂	iPr	11	-	-
PhCH ₂ CH ₂	iPr	11	-	-
2-naph-CH ₂	iPr	11	-	-
1-naph-CH ₂	iPr	11	-	-
4-thiazolyl- CH ₂	iPr	11	-	-

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R	R ¹	m	n	A
3-indolyl-CH ₂	iPr	11	-	-
PhCH ₂	iPr	-	10	O
PhCH ₂	iPr	-	12	O
PhCH ₂	iPr	2	2	S(CH ₂) ₃ S
PhCH ₂	iPr	2	2	S(CH ₂) ₄ S
PhCH ₂	iPr	2	2	S(CH ₂) ₅ S
PhCH ₂	iPr	2	2	S(CH ₂) ₆ S
PhCH ₂	iPr	2	2	S(CH ₂) ₇ S
PhCH ₂	iPr	2	2	SCH ₂ CH=CHCH ₂ S (cis)
PhCH ₂	iPr	2	2	SCH ₂ CH=CHCH ₂ S (trans)
PhCH ₂	iPr	2	2	SCH ₂ CCCH ₂ S
PhCH ₂	iPr	2	2	SCH ₂ p-Ph-CH ₂ S
PhCH ₂	iPr	2	2	SCH ₂ m-Ph-CH ₂ S
PhCH ₂	iPr	2	2	SCH ₂ o-Ph-CH ₂ S
PhCH ₂	iPr	2	2	SCH ₂ p-(Cl) ₄ Ph- CH ₂ S
PhCH ₂	iPr	11	-	replace amide NH by O.

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18. A compound of formula V:



where $X = \text{CWNR}, \text{S(O)}_n\text{NR}, \text{P(O)(Q)NR}$; $W = \text{O}, \text{S}, \text{NR}$; $n = 0, 1, 2$;

$Q = \text{R}, \text{OR}, \text{N(R)}_2$

$Y = \text{NRCW}, \text{NRS(O)}_n, \text{NRP(O)(Q)}$

$Z = \text{CRR}', \text{NR}$

A = an optionally substituted bridging group which is saturated or unsaturated and having 2-15 atoms comprised of any stable combination of C, N, O, or S.

R = represents hydrogen, alkoxy carbonyl, aralkoxy carbonyl, alkyl carbonyl, cycloalkyl carbonyl, cycloalkyl alkoxy carbonyl, cycloalkyl alkanoyl, aralkanoyl, aroyl, aryloxy carbonyl, aryloxy carbonyl alkyl, aryloxy alkanoyl, heterocyclyl carbonyl, heterocyclyloxy carbonyl, heterocyclyl alkanoyl, heterocyclyl alkoxy carbonyl, heteroaralkanoyl, heteroaralkoxy carbonyl, heteroaryloxy carbonyl, heteroaroyl, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, aryloxy alkyl, heteroaryloxy alkyl, hydroxy alkyl, aminocarbonyl, amino alkanoyl, mono- and disubstituted aminocarbonyl and mono- and di-substituted amino alkyl, alkoxy alkyl, alkylthio alkyl, mono- and disubstituted amino alkanoyl radicals wherein the substituents are selected from alkyl, aryl, aralkyl, cycloalkyl, cycloalkyl alkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heterocycloalkyl alkyl radicals, or where said amino alkanoyl radical is disubstituted, said substituents along with the nitrogen atom to which they are attached form a heterocycloalkyl or heteroaryl radical;

R', R_2 and $R_3 = \text{H}, \text{halo}, \text{optionally substituted lower alkyl}, \text{lower alkoxy}, \text{lower alkylthio}, \text{mono or di-lower alkyl amino}.$

19. The compound of claim 18, wherein $X, Y = \text{NRCW}, \text{NRS(O)}_n, \text{NRP(O)(Q)}$; $Z = \text{CRR}', \text{NR}, \text{O}, \text{S}.$

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20. The compound of claim 18 wherein X = OCWNR, SCWNR; Y = NRCW, NRS(O)_n, NRP(O)Q; Z = CRR', NR'.

21. The compounds of any one of claims 1, 5, 9, 13 or 18 in the form of an acid addition salt or complex thereof.

22. A pharmaceutical composition comprising at least one compound from any one of claims 1, 5, 9, 13 or 18 and a pharmaceutically acceptable carrier.

23. A method of inhibiting cathepsin D, plasmepsin I or plasmepsin II, which comprises contacting an environment containing cathepsin D, plasmepsin I or plasmepsin II with an inhibiting effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

24. A method of inhibiting the growth of a *Plasmodium* protozoan in a subject which comprises administering to said subject a growth inhibiting effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

25. The method of claim 24, wherein the *Plasmodium* protozoan is *Plasmodium falciparum*.

26. A method of treating malaria in a subject which comprises administering to said subject an effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

27. A method of inhibiting the invasion and metastasis of tumor cells in a subject which comprises administering to said subject an effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

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28. The method of claim 27, wherein the tumor cells are breast cancer cells.

29. A method of inhibiting the accumulation of amyloid β protein in a cell or a population of cells which comprises treating said cell or population of cells with an effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

30. A method of alleviating the effects of Alzheimer's disease in a subject having said disease comprising administering to said subject an effective amount of at least one compound from any one of claims 1, 5, 9, 13 or 18.

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/02930

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07K5/02 A61K38/05

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C07K A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	J.MED.CHEM., vol. 29, no. 12, 1986, USA, pages 2519-24, XP002034889 NIRANKAR S.AGARWAL ET AL.: "Inhibition of Cathepsin D by Substrate analogues Containing Statine and by Analogues of Pepstatin" see figure 1; tables I,II ---	1,3,5,7, 10,11, 21-23
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☒ Further documents are listed in the continuation of box C.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
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X	EP 0 401 676 A (BIO MEGA INC) 12 December 1990 see claim 7 ---	10,11, 21,22
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X	EP 0 443 573 A (BIO MEGA INC) 28 August 1991 see examples 3-10 ---	1,5,10, 21,22
X	EP 0 566 237 A (LILLY CO ELI) 20 October 1993 see claim 1 ---	1,5,10, 21,22
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X	NAT.PROD.LETT., vol. 5, no. 2, 1994, pages 89-93, XP002034892 H.WU ET AL.: "The Structure of Qingjingmycin a Novel Peptide-Type Metabolite of a Hybrid Strain FM3-32 Derived from Streptomyces Qingfengmyceticus and S.Hydroscopius Var. Jinggangensis" see figure 1 ---	13,18, 21,22
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